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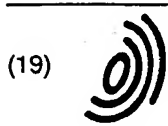
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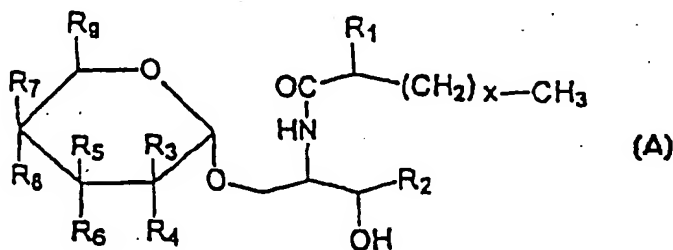
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(54) METHOD FOR ACTIVATING HUMAN ANTIGEN PRESENTING CELLS, ACTIVATED HUMAN ANTIGEN PRESENTING CELLS, AND USE OF THE SAME

(57) A method for activating human antigen presenting cells characterized by culturing *in vitro* antigen presenting cells with a human origin together with at least one of glycoside compounds represented by general formula (A) or salts thereof preferably exemplified by (2S, 3S, 4R)-1-(α -D-galactopyranosyloxy)-2-hexacosanoylamino-3,4-octadecanediol; the human antigen presenting cells thus activated; methods for treating cancer or infectious diseases such as AIDS by using activated human antigen presenting cells; and the use of activated human antigen presenting cells in the production of drugs for treating these diseases. Thus, sufficient therapeutic effects can be achieved on cancer and infectious diseases such as AIDS without effecting any pulse therapy with the use of tumor antigens.



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Description

BACKGROUND OF THE INVENTION

5 1. Field of the Invention

[0001] The present invention relates to a novel activated human antigen-presenting cell, a method for activating a human antigen-presenting cell *in vitro*, a method for treating cancer and infectious diseases including AIDS using the activated human antigen-presenting cell, and a use of the activated human antigen-presenting cell in preparing medicines for such treatment.

2. Disclosure of Related Art

[0002] There have been various strategies for the therapeutical treatment of cancer patients, including surgical extraction of tumor foci, chemotherapy, radiotherapy and immunotherapy. However, the cure rates of these strategies are not as high as they are expected. Therefore, there exists a strong demand for developing new therapeutic agents and methods which can provide improved cure rates in the treatment of cancer. Macrophages, B cells and dendritic cells, which are members of antigen-presenting cell (APC), have been known to be cells essential for immune response. Recently, an idea that using APCs, particularly dendritic cells, to induce cancer immunity may be effective in treating cancer (Grabbe, S. et al., 1995, Immunol. Today, 16, 117) has attracted much attention.

[0003] Several processes for preparing dendritic cells from mouse have been known, such as from the spleen (Clowley, M. et al., 1989, Cell. Immunol., 118, 108), from the bone marrow (Inaba, K. et al., 1992, J. Exp. Med., 176, 1693) and from the epidermis (the epidermis-derived dendritic cells are known as "Langerhans's cells") (Witmer-Pack, M. et al., 1987, J. Exp. Med., 166, 1487). Furthermore, it was demonstrated that pulsing the dendritic cells prepared from the murine bone marrow with tumor antigen and administering the pulsed dendritic cells into a subject prior to and after the implantation of tumor cells into the subject can elicit tumor immunity (Celluzzi, C. M. et al., 1996, J. Exp. Med., 183, 283; Paglia, P. et al., 1996, J. Exp. Med., 183, 317).

[0004] The following processes are known for preparing human APCs, particularly dendritic cells, from peripheral blood or umbilical cord blood: a process in which FcR⁺ cells, T cells, B cells and NK cells are removed from human peripheral blood using the antibodies against them from peripheral blood monocytes to give APCs (Hsu et al., 1996, Nature Med., 2, 52); and a process in which adherent cells in human peripheral blood monocytes from which CD19⁺ B cells and CD2⁺ T cells were removed are cultured with a granulocyte-macrophage colony stimulating factor (GM-CSF) and interleukin 4 (IL-4) for about one week (Sallusto, F. et al., 1994, J. Exp. Med., 179, 1109). It has been reported that such human peripheral blood-derived APCs have both allogeneic and autologous MLR (mixed leukocyte reaction) enhancing effects. On the other hand, for preparing APCs from human umbilical cord blood or bone marrow cells, a process is known where GM-CSF and tumor necrosis factor- α (TNF- α) are used to prepare APCs from CD34⁺ cells found in umbilical cord blood or bone marrow (Santiago-Schwartz, F. et al., 1992, J. Leukocyte Biol., 52, 274; Caux, C. et al., 1992, Nature, 360, 258). However, although the APCs prepared from human umbilical cord blood or bone marrow cells by such a process as mentioned above have an allogeneic MLR enhancing effect, they have no (Santiago-Schwartz, F. et al.) or, if any, extremely poor (Caux et al.) autologous MLR enhancing effect.

[0005] Recently, it has been demonstrated that a therapy with the APCs that had been pulsed with tumor antigens is effective for treating B cell lymphoma patients (Hsu, F. J. et al., 1996, Nature Med., 2, 52). That is, a therapy for B cell lymphoma patients was successfully achieved by culturing APCs prepared from peripheral blood of a B cell lymphoma patient together with B cell lymphoma antigens *in vitro* and administering the cultured APCs into the B cell lymphoma patient.

[0006] As shown in this exemplary therapy, methods in which APCs are pulsed with tumor antigens may be very effective for treating cancer if the cancer can be clearly identified by their tumor antigens, like B cell lymphoma. However, such methods as mentioned above take enormous time and costs, because the tumor antigens are generally specific to the individual patients and, therefore, must be specified for the respective patients, and the specified tumor antigens must be produced in large quantities to pulse the APCs. Furthermore, since it is impossible for most of the cancer patients to identify their tumor antigens, the range of application of the method in which APCs are pulsed with tumor antigens is limited. In view of the above, in order to apply the therapy with APCs (hereinafter referred to as "the APC therapy") recognized to be a very effective therapy for cancer, to as many patients as possible, there has been a strong demand for development of a method for preparing human APCs effective for treating cancer without the aid of tumor antigens.

[0007] Within a living body, various types of β -galactosylceramides and β -glucosylceramides are present, each of which has a sugar linked to a ceramide via a β -linkage (Svennerholm, L. et al., 1972, Biochem. Biophys. Acta., 280, 626; Karlsson, K. A. et al., 1973, Biochim. Biophys. Acta., 316, 317). On the other hand, the inventors have found that

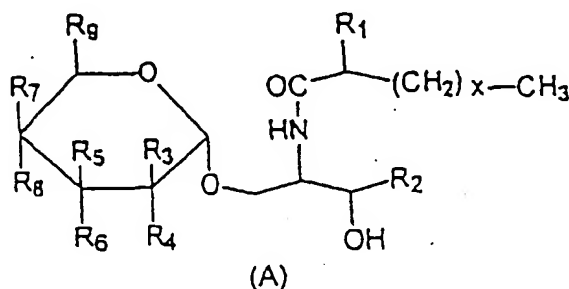
α -galactosylceramides have remarkable immunostimulatory properties and anti-tumor properties (Morita, M. et al., 1995, J. Med. Chem., 38, 2176), and that such properties of α -galactosylceramides and α -glucosylceramides are far more potent than those of the corresponding β -anomers (Motoki, K. et al., 1995, Biol. Pharm. Bull., 18, 1487). It was also found that compounds having an α -glycosylceramide structure show a protecting effect against radiation when administered into a living subject (Motoki, K. et al., 1995, Bioorg. Med. Chem. Lett., 5, 2413) and cause an increase in the number of platelets and white blood cells (Motoki, K. et al., 1996, Biol. Pharm. Bull., 19, 952). The inventors also have found that an α -galactosylceramide, KRN7000, can enhance the functions of dendritic cell-rich APCs prepared from the murine spleen, and that the enhanced APCs exhibit an anti-tumor effect when administered into a subject before tumors are implanted into the subject (Koezuka, Y. et al., 1995, The 9th International Congress of Immunology, Abstract, 55; Yamaguchi, Y. et al., 1996, Oncol. Res. 8, 399).

[0008] However, the effect of glycoside compounds, such as KRN7000, or salts thereof on APCs rich in dendritic cells from murine bone marrows and Langerhans's cells from murine epidermis has not been clarified. In addition, there is no report on anti-tumor effects of the APCs rich in murine spleen- and bone marrow-derived dendritic cells, which have been stimulated with KRN7000, upon administration into a subject after tumors are implanted into the subject. No effect of glycoside compounds such as KRN7000 or salts thereof on human APCs has also been known.

[0009] To address the above-mentioned demands, the present invention provides a novel human APC activating agent which is useful in preparing activated human APCs that exhibit a sufficient therapeutic effect on cancer and various infectious diseases including AIDS without the need to pulse the antigen-presenting cells with tumor antigens, and also provides a method for activating human APCs with such an activating agent.

SUMMARY OF THE INVENTION

[0010] The present invention provides a method for activating human APCs, which comprises culturing human-derived APCs *in vitro* with at least one of glycoside compounds represented by formula (A) or salts thereof:



wherein R_1 to R_9 and X are to be defined later.

[0011] According to one embodiment, the present invention provides a method for activating human APCs which comprises culturing human-derived APCs *in vitro* with a glycoside compound (2S, 3S, 4R)-1-(α -D-galactopyranosyloxy)-2-hexacosanoylamino-3,4-octadecanediol or a salt thereof.

[0012] The present invention also provides activated human APCs which can be prepared by culturing human-derived APCs *in vitro* with at least one of the glycoside compounds represented by formula (A) above or salts thereof.

[0013] The present invention further provides a method for treating cancer and infectious diseases including AIDS, which comprises applying an APC therapy with such activated human APCs as defined above.

[0014] The present invention further provides a use of such activated human APCs as defined above in the preparation of a medicine for treating cancer and infectious diseases including AIDS by application of the APC therapy.

BRIEF DESCRIPTION OF THE DRAWINGS

[0015]

Fig. 1 shows the amount of ^3H -TdR taken up into the cells variously treated as described. In Fig. 1, "V-APC" and "KRN-APC" refer to human peripheral blood-derived APCs pretreated with a vehicle and KRN7000, respectively. Fig. 2 shows a relationship between the number of APCs and the amount of ^3H -TdR taken up into an allogeneic T cells. In Fig. 2, "V-APC" and "KRN-APC" refer to human umbilical cord blood-derived APCs (CD1c^+ cells) stimulated with a vehicle and KRN7000, respectively.

Fig. 3 shows a relationship between the number of APCs and the amount of ^3H -TdR taken up into autologous T

cells. In Fig. 3, "V-APC" and "KRN-APC" refer to the same APCs as those used in Fig. 2.

Fig. 4 shows a relationship between the number of APCs and the amount of ^3H -TdR taken up into spleen cells. In Fig. 2, "V-APC", "KRN-APC", "583-APC", "517-APC", "564-APC", "563-APC" and "562-APC" refer to murine spleen-derived APCs pretreated with a vehicle, KRN7000, AGL-583, AGL-517, AGL-564, AGL-563 and AGL-562, respectively.

Fig. 5 shows the survival periods of EL-4-bearing mice when murine spleen-derived APCs pretreated with a vehicle or KRN7000 were injected intravenously to the mice.

Fig. 6 shows a relationship between the number of APCs and the amount of ^3H -TdR taken up into T cells. In Fig. 6, "V-APC" and "KRN-APC" refer to murine bone marrow-derived APCs pretreated with a vehicle and KRN7000, respectively.

Fig. 7 shows the weight of the liver of a tumor-bearing mouse when the V-APC or KRN-APC prepared in the same manner as in Fig. 6 was injected intravenously into the mouse.

Fig. 8 shows a relationship between the number of APCs and the amount of ^3H -TdR taken up into spleen cells. In Fig. 8, "V-APC" and "KRN-APC" refer to murine epidermis-derived APCs pretreated with a vehicle and KRN7000, respectively.

Fig. 9 shows the volume of a subcutaneous tumor for each of the tumor-bearing mice when the APCs pretreated with a vehicle, a vehicle+B16-tumor cell lysate, KRN7000 or KRN7000+B16-tumor cell lysate (V-APC, V-T-APC, KRN-APC or KRN-T-APC, respectively), respectively, was injected intravenously to the mice.

Fig. 10 shows a relationship between the number of APCs given after culturing for 3 days and the amount of ^3H -TdR taken up into human peripheral blood-derived FcR $^+$ cells. In Fig. 10, "V-APC", "KRN-APC", "V-APC-GM" and "KRN-APC-GM" refer to human peripheral blood-derived APCs pretreated with a vehicle, KRN7000, a vehicle+GM-CSF+IL4 and KRN+GM-CSF+IL4, respectively.

Fig. 11 shows a relationship between the number of APCs given after culturing for 7 days and the amount of ^3H -TdR taken up into human peripheral blood-derived FcR $^+$ cells. In Fig. 11, "V-APC-GM" and "KRN-APC-GM" refer to the same APCs as those used in Fig. 10.

Fig. 12 shows a relationship between the number of APCs given after culturing for 11 days and the amount of ^3H -TdR taken up into human peripheral blood-derived FcR $^+$ cells. In Fig. 12, "V-APC-GM" and "KRN-APC-GM" refer to the same APCs as those used in Fig. 10, and "MCM $^+$ " and "MCM $^-$ " respectively mean that the APCs are cultured with and without a monocyte conditioned medium (MCM).

Fig. 13 shows a relationship between the number of APCs given after culturing for 3 days and the amount of ^3H -TdR taken up into human umbilical cord blood-derived FcR $^+$ cells. In Fig. 13, "V-APC-GM" and "KRN-APC-GM" refer to the human umbilical cord blood-derived APCs respectively pretreated with a vehicle+GM-CSF+IL4 and KRN7000+GM-CSF+IL4.

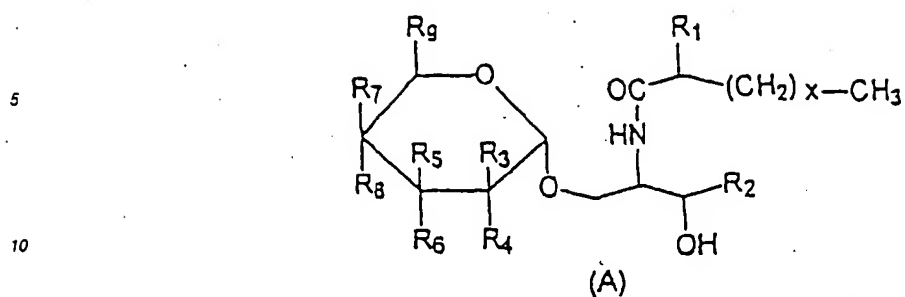
Fig. 14 shows activation effect of murine spleen-derived APCs when they are stimulated by Compounds Nos. 1, 2, 5, 10 and 16 (except KRN7000), termed AGL506, AGL514, AGL571, AGL512 and AGL525, respectively.

DETAILED DESCRIPTION OF THE INVENTION

[0016] As a result of extensive and intensive studies by the inventors on the above-mentioned problems, it was found that the glycoside compounds according to the present invention (representatively, KRN7000) can significantly enhance the functions of human APCs prepared from human peripheral blood and human umbilical cord blood. The glycoside compounds can also significantly enhance the functions of dendritic cell-rich APCs from a murine bone marrow and Langerhans's cell-rich APCs from murine auricle, as well as dendritic cell-rich APCs from a murine spleen. It was also found that a significant anti-tumor effect can be observed when dendritic cell-rich APCs from a murine spleen or a murine bone marrow which have been cultured *in vitro* with the glycoside compounds of the present invention (representatively, KRN7000) are injected into a mouse into which tumor cells have been implanted. Based on these findings, the invention was completed.

[0017] That is, the present invention relates to an activating agent (i.e., an antigen-presenting function enhancing agent) comprising a specific glycoside compound, which is useful for preparing activated human APCs having a therapeutic effect on cancer and infectious diseases including AIDS. Specifically, the present invention relates to an activating agent for APCs prepared from human peripheral blood, human umbilical cord blood, human bone marrow cells or human epidermis (i.e., APCs to be activated). More specifically, the present invention relates to an activating agent for human monocytes, human dendritic cells or human CD1c $^+$ cells.

[0018] According to the present invention, there is provided a method for activating human antigen-presenting cells which comprises culturing human-derived antigen-presenting cells *in vitro* with at least one of the glycoside compounds represented by formula (A) or salts thereof:



15 wherein:

R_1 is H or OH;

X is an integer of from 7 to 25;

R_2 is a substituent defined by any one of the following (a) to (e):

20

(a) $-\text{CH}_2(\text{CH}_2)_Y\text{CH}_3$;

(b) $-\text{CH}(\text{OH})(\text{CH}_2)_Y\text{CH}_3$;

(c) $-\text{CH}(\text{OH})(\text{CH}_2)_Y\text{CH}(\text{CH}_3)_2$;

(d) $-\text{CH}=\text{CH}(\text{CH}_2)_Y\text{CH}_3$; and

25

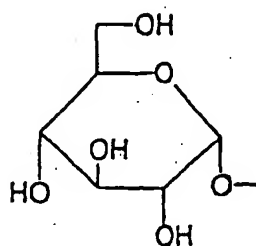
(e) $-\text{CH}(\text{OH})(\text{CH}_2)_Y\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_3$;

wherein Y is an integer of from 5 to 17;

R_3 is H;

R_4 is OH, NH_2 , NHCOCH_3 or

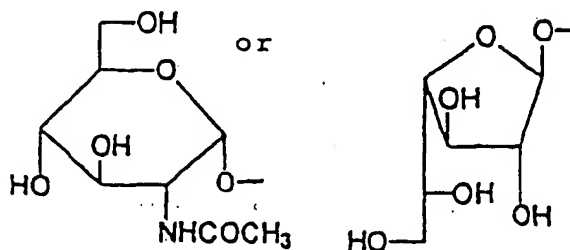
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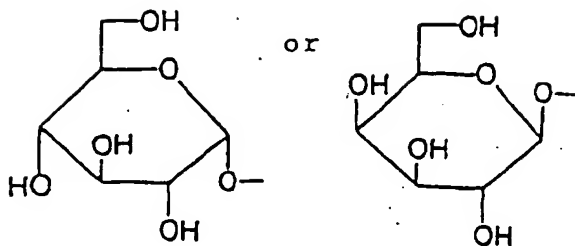
one of R_5 and R_6 is H and the other is OH,

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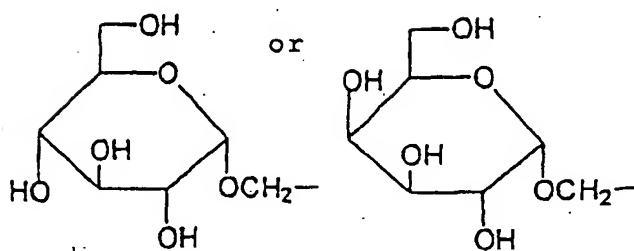


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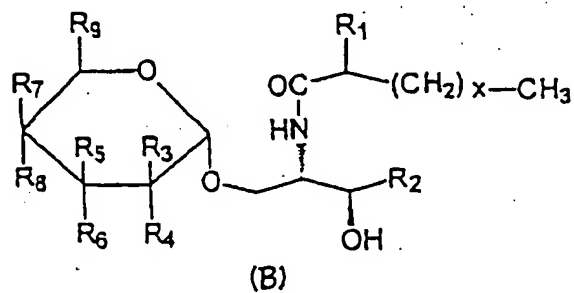
one of R_7 and R_8 is H and the other is OH,



and
 R_9 is H, CH_3 , CH_2OH ,



[0019] In a preferred embodiment, the present invention provide a method for activating human antigen-presenting cells which comprises using at least one of the glycoside compounds represented by formula (B) or salts thereof:



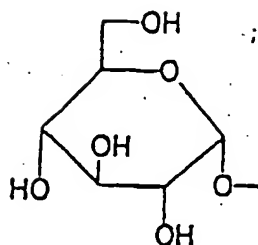
wherein:

R_1 , X and R_2 are as defined as in the case of formula (A); and
 R_3 to R_9 are substituents defined by any one of the following (i) to (iii):

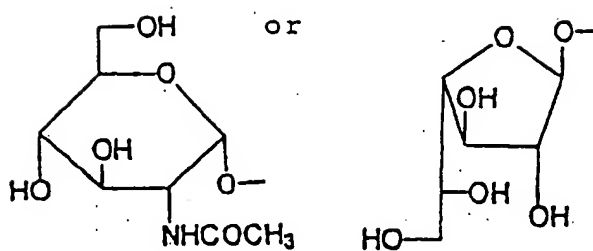
(i) [galactose type]

each of R_3 , R_6 and R_8 is H;

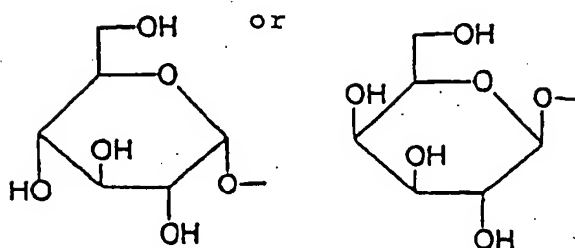
R_4 is OH or



15 R_5 is OH,

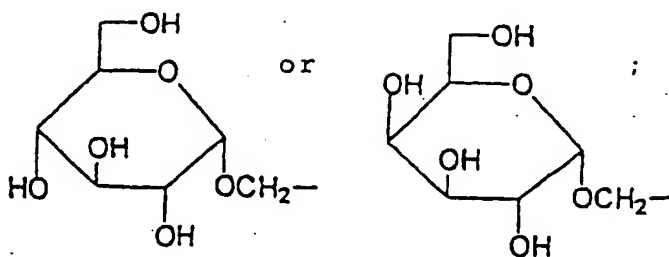


30 R_7 is OH,



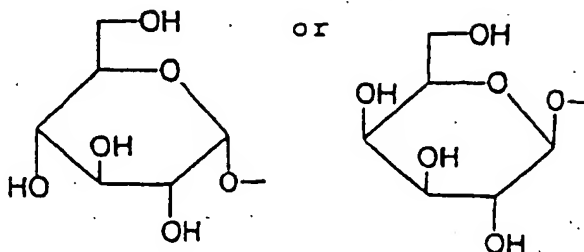
and

45 R_9 is H, CH_3 , CH_2OH ,



(ii) [glucose type]

each of R_3 , R_6 and R_7 is H;
 R_4 , R_5 and R_9 are as defined as in (i); and
 R_8 is OH,



or

(iii) [allose type]

each of R_3 , R_5 and R_7 is H;
each of R_4 , R_6 and R_8 is OH; and
 R_9 is H, CH_3 or CH_2OH .

[0020] The glycoside compounds defined by formula (A) or (B) above are comprised of a sugar moiety and an aglycone moiety, and some of them are also referred to as α -cerebrosides, α -glycosylceramides, α -glucosylceramides, α -galactocerebrosides or α -galactosylceramides. These compounds are characterized by having the α -form of anomeric configuration.

[0021] In the glycoside compound, the sugar moiety is preferably of [galactose type] as defined in (i), and more preferably of one wherein each of R_3 , R_6 and R_8 is H, each of R_4 , R_5 and R_7 is OH and R_9 is CH_2OH (i.e., α -galactopyranosyl).

[0022] In the glycoside compound, the aglycone moiety preferably has R_2 being any one of the substituents (b), (c) and (e) above, and more preferably has R_1 being H (i.e., kerosin type) and R_2 being the substituent (b). X is preferably an integer of 21 to 25 and Y is preferably an integer of 11 to 15.

[0023] Preferable examples of the glycoside compound of the present invention are listed below. In the list, compounds (1)-(9), (10)-(24), (25)-(31), (32)-(33), and (34) are those compounds in which R_2 is the substituent (a), (b), (c), (d) or (e) above, respectively. The alphabet letters A, B, C and D behind the compounds' name indicate the reference specifications of WO93/05055, WO94/02168, WO94/09020 and WO94/24142, respectively, which describe the synthesis methods of the annotated compounds. Among the glycoside compounds below, compound (14) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-hexacosanoylamino-3,4-octadecanediol (referred to as "KRN7000" hereinbelow), is most preferable. With respect to this compound, an example of the synthesis process will be illustrated in the Production Example and Scheme 1 below.

(1) (2S,3R)-1-(α -D-galactopyranosyloxy)-2-[(R)-2-hydroxytetracosanoylamino]-3-octadecanol	A
(2) (2S, 3R)-1-(α -D-galactopyranosyloxy)-2-tetracosanoylamino]-3-octadecanol	A
(3) (2S,3R)-1-(α -D-galactopyranosyloxy)-2-tetradecanoylamino-3-octadecanol	A
(4) (2S,3R)-1-(α -D-glucopyranosyloxy)-2-tetradecanoylamino-3-octadecanol	C
(5) (2S,3R)-1-(6'-deoxy- α -D-galactopyranosyloxy)-2-tetradecanoylamino-3-octadecanol	C
(6) (2S,3R)-1-(β -L-arabinopyranosyloxy)-2-tetradecanoylamino-3-octadecanol	C
(7) (2S,3R)-1-(α -D-galactopyranosyloxy)-2-tetradecanoylamino-3-hexadecanol	A

(continued)

	(8) (2R,3R)-1-(α -D-galactopyranosyloxy)-2-tetradecanoylamino-3-hexadecanol	A
	(9) (2R,3S)-1-(α -D-galactopyranosyloxy)-2-tetradecanoylamino-3-hexadecanol	A
5	(10) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(R)-2-hydroxytetraacosanoylamino]-3,4-octadecanediol	A
	(11) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(R)-2-hydroxytetraacosanoylamino]-3,4-undecanediol	A
10	(12) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(R)-2-hydroxyhexacosanoylamino]-3,4-icosanediol	A
	(13) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(S)-2-hydroxytetraacosanoylamino]-3,4-heptadecanediol	A
15	(14) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-hexacosanoylamino-3,4-octadecanediol	Production Example
	(15) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-octacosanoylamino-3,4-heptadecanediol	B
	(16) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-tetracosanoylamino-3,4-octadecanediol	A
20	(17) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-tetracosanoylamino-3,4-undecanediol	A
	(18) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-hexacosanoylamino-3,4-octadecanediol	C
	(19) O- β -D-galactofuranosyl-(1 \rightarrow 3)-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxytetraacosanoyl]-1,3,4-octadecanetriol	D
25	(20) O- α -D-galactopyranosyl-(1 \rightarrow 6)-O- α -D-glucopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-hexacosanoyl-1,3,4-octadecanetriol	D
	(21) O- α -D-galactopyranosyl-(1 \rightarrow 6)-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-hexacosanoyl-1,3,4-octadecanetriol	D
30	(22) O- α -D-glucopyranosyl-(1 \rightarrow 4)-O- α -D-glucopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-hexacosanoyl-1,3,4-octadecanetriol	D
	(23) O-(N-acetyl-2-amino-2-deoxy- α -D-galactopyranosyl-(1 \rightarrow 3))-O-[α -D-glucopyranosyl-(1 \rightarrow 2)]-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxyhexacosanoyl]-1,3,4-octadecanetriol	D
35	(24) O-(N-acetyl-2-amino-2-deoxy- α -D-galactopyranosyl-(1 \rightarrow 3))-O-[α -D-glucopyranosyl-(1 \rightarrow 2)]-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxytetraacosanoyl]-1,3,4-hexadecanediol	D
40	(25) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(R)-2-hydroxytricosanoylamino]-16-methyl-3,4-heptadecanediol	A
	(26) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(S)-2-hydroxytetraacosanoylamino]-16-methyl-3,4-heptadecanediol	A
45	(27) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-16-methyl-2-tetracosanoylamino]-3,4-heptadecanediol	A
	(28) O- β -D-galactofuranosyl-(1 \rightarrow 3)-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxytetraacosanoyl]-17-methyl-1,3,4-octadecanetriol	D
50	(29) O- β -D-galactofuranosyl-(1 \rightarrow 3)-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxytetraacosanoyl]-15-methyl-1,3,4-hexadecanediol	D
	(30) O-(N-acetyl-2-amino-2-deoxy- α -D-galactopyranosyl-(1 \rightarrow 3))-O-[α -D-glucopyranosyl-(1 \rightarrow 2)]-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxyhexacosanoyl]-16-methyl-1,3,4-octadecanetriol	D
55	(31) O-(N-acetyl-2-amino-2-deoxy- α -D-galactopyranosyl-(1 \rightarrow 3))-O-[α -D-glucopyranosyl-(1 \rightarrow 2)]-O- α -D-galactopyranosyl-(1 \rightarrow 1)-(2S,3S,4R)-2-amino-N-[(R)-2-hydroxytetraacosanoyl]-16-methyl-1,3,4-heptadecanetriol	D

(continued)

	(32) (2S,3S,4E)-1-(α -D-galactopyranosyloxy)-2-octadecanoylamino-4-octadecene-3-ol	A
	(33) (2S,3S,4E)-1-(α -D-galactopyranosyloxy)-2-tetradecanoylamino-4-octadecene-3-ol	A
5	(34) (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-[(R)-2-hydroxypentacosanoylamino]-16-methyl-3,4-octadecanediol	A

10 The glycoside compound defined by formula (A) or (B) may form an acid addition salt with a pharmaceutically acceptable acid. Examples of the acid to be used for formation of such an acid addition salt include inorganic acids such as hydrochloric acid, sulfuric acid, nitric acid and phosphoric acid; and organic acids such as acetic acid, propionic acid, maleic acid, oleic acid, palmitic acid, citric acid, succinic acid, tartaric acid, fumaric acid, glutamic acid, pantothenic acid, lauryl sulfonic acid, methanesulfonic acid and phthalic acid.

15 [0024] The human APC-activating agent according to the present invention can be used to activate the APCs prepared from various human tissues such as peripheral blood, umbilical cord blood, bone marrow cells and epidermis. Typically, the activating agent of the present invention can be added *in vitro* during or after the preparation of the human APCs (for example, from human peripheral blood, umbilical cord blood or bone marrow cells and, if necessary, with GM-CSF and IL-4) to give the activated human APCs. The activated human APCs may be prepared more efficiently by adding substance(s) such as SCF, IL-1, IL-3, TNF- α and Flk2/Flt3 ligand or a monocyte conditioned medium (MCM) in addition to GM-CSF and IL-4. Moreover, in the preparation of the activated human APCs, tumor antigens, viral antigens or antigen-specific peptides thereof can be added to give the activated human APCs that are more effective for treating cancer and infectious diseases.

20 [0025] The human APC-activating agent of the present invention may be dissolved in an appropriate dissolving solvent such as DMSO and then diluted to an appropriate concentration with physiological saline, a culture medium or the like. The final concentration of the activating agent in a suitable cultivation for activating human APCs is within the range from 0.1 ng/mL to 10 μ g/mL, and preferably from 0.01 μ g/mL to 1 μ g/mL. It is preferable that the human APC-activating agent according to the present invention be prepared and added immediately before it is used.

25 [0026] Accordingly, the present invention also provides activated human APCs which can be prepared by culturing human-derived APCs *in vitro* with at least one of the glycoside compounds represented by formula (A) or (B) defined above or salts thereof.

30 [0027] In an embodiment of the present invention, the human-derived APCs are CD1c-positive or CD1c-rich cells. Such cells may be prepared from, but not limited to, human peripheral blood, human umbilical cord blood, human bone marrow cells, human epidermis, and the like.

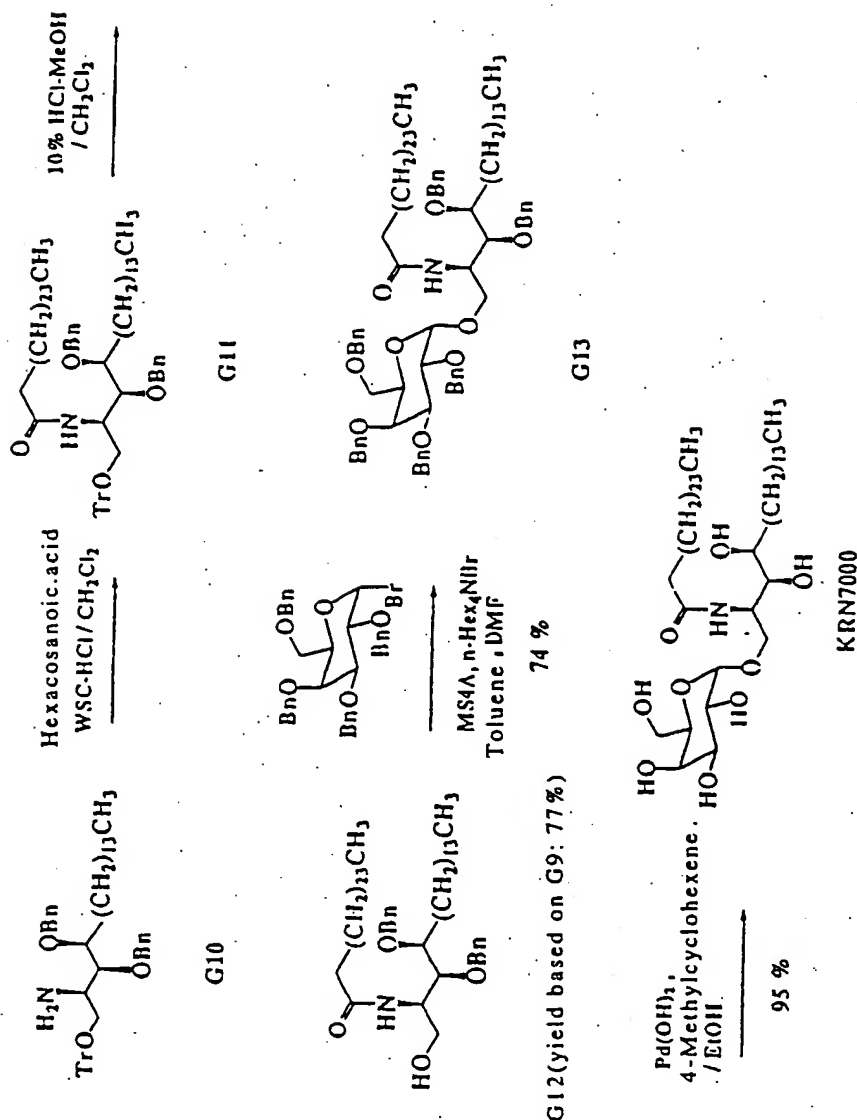
35 [0028] The use of the human APCs activated with the activating agent of the present invention leads to effective APC therapy for patients suffering from cancer. Furthermore, it is expected that the cytotoxic T lymphocytes (CTLs) prepared by co-culturing the activated APCs and peripheral blood from a patient can be used for an effective therapy for patients suffering from cancer or infectious diseases including AIDS.

[0029] Accordingly, the present invention also provides a method for treating cancer and infectious diseases including AIDS with the activated human APCs.

40 [0030] The present invention further provides a use of the activated human APCs in the preparation of medicines for treating cancer and infectious diseases including AIDS by application of an APC therapy.

[0031] The following examples illustrate the present invention more clearly; however, these examples are not be construed to limit the scope of the invention.

SCHEME 1 (continued)

45 Synthesis of Compound G1

[0033] To a solution of D-lyxose (200 g, 1.33 mol) in acetone (3.0 L) dried over calcium chloride was added sulfuric acid (0.5 mL). The mixture was stirred at room temperature for 18 hours, and Molecular Sieves 4A powder (100 g) was then added thereto. After neutralizing, the reaction solution was filtered with celite and the residue was washed with acetone. The filtrate and the washings were combined and concentrated under reduced pressure, thereby giving a crude product of Compound G1 in an amount of 240 g (yield 95%). For the subsequent reaction, the crude product was used without any further purification. For the preparation of a sample for analyses, a portion of the crude product was subjected to further purification by chromatography on a silica gel column with hexane:acetone (9:1) as an eluent.

55 mp 76-78°C; FDMS m/z 191 ($M+1$)⁺; ¹H-NMR (500MHz, CDCl₃) δ 5.45 (1H, d, J = 1.8Hz), 4.83 (1H, dd, J = 3.7, 5.5Hz), 4.64 (1H, d, J = 6.1Hz), 4.27-4.30 (1H, m), 3.90-3.99 (2H, m), 1.48 (3H, s), 1.32 (3H, s).

Synthesis of Compound G2

[0034] To a solution (168 mL) of Compound G1 (239 g, about 1.26 mol) in methylene chloride were added pyridine (10 mL) and trityl chloride (39.0 g), and the mixture was stirred at 32°C for 4 hours. Ethanol (8 mL) was then added to the solution dropwise and stirred at room temperature for 2 hours. The solution was washed with a saturated aqueous ammonium chloride solution, a saturated aqueous sodium hydrogencarbonate solution and the brine, and was concentrated under reduced pressure. The residue was dissolved in ethyl acetate and then cooled to 0°C to cause precipitation, thereby giving a product of Compound G2 in an amount of 501 g (yield based on the amount of D-lyxose: 87%).

mp 174-176°C; FDMS m/z 432 M⁺; ¹H-NMR (500MHz, CDCl₃) δ 7.21-7.49 (15H, m), 5.38(1H, d, J = 2.4Hz), 4.75(1H, dd, J = 3.7, 6.1Hz), 4.59(1H, d, J = 6.1Hz), 4.31-4.35(1H, m), 3.43 (1H, dd, J = 4.9, 9.8Hz), 3.39(1H, dd, J = 6.7, 9.8Hz), 1.29(3H,s), 1.28(3H,s).

Synthesis of Compound G3

[0035] To a solution (1500 mL) of tridecanetriphenylphosphonium bromide (962 g, 1.16 mol; a product prepared by heating 1-bromotridecane and triphenylphosphine at 140°C for 4.5 hours) in THF was added a 2.5 M solution of n-butyl lithium in hexane (462 mL, 1.16 mol) dropwise in an atmosphere of argon at 0°C. After the dropwise addition was completed, the resultant solution was stirred for 15 minutes. To the solution was added a solution (450 mL) of Compound G2 (250 g, 579 mmol) in THF dropwise. The resultant solution was stirred for 18 hours while gradually elevating the solution temperature to room temperature. The reaction solution was concentrated under reduced pressure. To the residue was added a mixed solution of hexane:methanol:water (10:7:3, 1000 mL) and washed with a saturated aqueous ammonium chloride solution. The aqueous layer was extracted with hexane (500 mL) several times. All of the organic layers obtained were combined, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure, thereby giving a crude product of Compound G3 in an amount of 339 g (yield: 98%). For the subsequent reaction, the crude product was used without any further purification. For the preparation of a sample for analyses, a portion of the crude product was subjected to further purification by chromatography on a silica gel column with hexane:ethyl acetate (9:1) as an eluent.

FDMS m/z 598M⁺; ¹H-NMR (500MHz, CDCl₃) δ 7.21-7.45 (15H, m), 5.48-5.59(2H, m), 4.91(0.7H, t, J = 7.3Hz), 4.44(0.3H, t, J = 7.3Hz), 4.26(0.3H, dd, J = 4.3, 7.3Hz), 4.21(0.7H, dd, J = 4.3, 6.7Hz), 3.75(0.7H, m), 3.69(0.3H, m), 3.24(0.3H, dd, J = 4.9, 9.8Hz), 3.17(0.7H, dd, J = 4.9, 9.8Hz), 3.09-3.14 [1H, (3.11, dd, J = 4.9, 9.2Hz), H1bE overlapped], 1.75-2.03(2H, m), 1.49(3H, s), 1.39 and 1.38(3H, each s), 1.21-1.34(20H, m), 0.88(3H, t, J = 6.7Hz).

Synthesis of Compound G4

[0036] To a solution (1500 mL) of Compound G3 (338 g, about 565 mmol) in methylene chloride was added pyridine (500 mL). Subsequently methanesulfonyl chloride (49 mL, 633 mmol) was added dropwise thereto and then stirred at 31°C for 24 hours. Ethanol (40 mL) was added dropwise thereto and stirred at room temperature for 1 hour. After the reaction solution was concentrated under reduced pressure, a mixed solution of hexane:methanol:water (10:7:3, 1000 mL) was added to the residue to cause separation of the solution into an aqueous layer and an organic layer. The aqueous layer was extracted with hexane (200 mL) 3 times. All of the organic layers obtained were combined, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure, thereby giving a crude product of Compound G4 in an amount of 363 g (yield: 95%). For the subsequent reaction, the crude product was used without any further purification. For the preparation of a sample for analyses, a portion of the crude product was subjected to further purification by chromatography on a silica gel column with hexane:ethyl acetate (9:1) as an eluent.

FDMS m/z 676M⁺; ¹H-NMR (500MHz, CDCl₃) δ 7.21-7.47 (15H, m), 5.41(0.7H, ddd, J = 5.5, 9.2, 11.0 Hz), 5.32(0.7H, bt, J = 11.0 Hz), 5.22 (0.3H, bdd, J = 9.2, 15.0 Hz), 5.02 (0.3H, dt, J_t = 7.3Hz, J_d = 15.0 Hz), 4.8 (0.7H, ddd, J = 3.1, 5.5, 7.9Hz), 4.73(0.7H, dd, J = 5.5, 9.8 Hz), 4.64-4.67 (0.3H, m), 4.61(0.3H, dd, J = 5.5, 9.2Hz), 4.48(0.7, dd, J = 5.5, 7.9Hz), 4.22(0.3H, dd, J = 5.5, 9.2 Hz), 3.55(0.3H, dd, J = 2.4, 11.6 Hz), 3.45(0.7H, dd, J = 3.2, 11.0 Hz), 3.06-3.12[4H, (3.12,S), (3.11,S), (3.09, dd, J = 3.1, 11.0 Hz)], 1.66-1.82 (2H,m), 1.47 and 1.46 (3H, each s), 1.39 (3H,s), 1.13-1.35 (20H,m), 0.88 (3H, t, J = 6.8 Hz).

Synthesis of Compound G5

[0037] To a solution (1500 mL) of Compound G4 (362 g, about 536 mmol) in methylene chloride was added methanol (350 mL). Concentrated hydrochloric acid (200 mL) was added to the solution dropwise and then stirred at room tem-

perature for 5 hours. The reaction solution was neutralized with sodium hydrogencarbonate and then subjected to filtration. The filtrate was concentrated under reduced pressure, and ethyl acetate was added to the residue and then washed with brine. The aqueous layer was extracted with ethyl acetate several times. All of the organic layers obtained were combined, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure. The resultant product was crystallized from hexane to give a product of Compound G5 in an amount of 161 g (yield based on the amount of Compound G2: 70%).

mp 66-67°C ; FDMS m/z 377(M- H₂O)+ ; ¹H-NMR(500MHz, CDCl₃+D₂O) δ 5.86(0.3H, dt, J_t = 7.3Hz, J_d = 14.7Hz), 5.77(0.7H, dt, J_t = 7.3, J_d = 10.4Hz), 5.55(0.3H, br.dd, J = 7.3,14.7 Hz), 5.49(0.7H, bt, J = 9.8 Hz), 4.91-4.97(1H, m), 4.51 (0.7H, bt, J = 9.8Hz), 4.11(0.3H, bt, J = 7.3Hz), 3.94-4.03 (2H,m), 3.67-3.73 [1H, (3.70, dd, J = 3.1, 6.7Hz), (3.69, dd, J = 3.1, 7.3 Hz)], 3.20 and 3.19 (3H,each s), 2.05-2.22(2H,m), 1.22-1.43(20H,m), 0.88(3H, t, J = 6.7Hz).

Synthesis of Compound G6

[0038] To a solution (780 mL) of Compound G5 (160 g, 405 mmol) in THF was added 5% palladium on barium sulfate (16 g). After the reaction vessel was purged with hydrogen gas, the mixture was stirred at room temperature for 20 hours. The reaction solution was filtered with celite and the filter cake was washed with a mixed solution of chloroform:methanol (1:1). The filtrate and the washings were combined and concentrated under reduced pressure. The residue was crystallized from ethyl acetate to give a product of Compound G6 in an amount of 146 g (yield: 91%).

[α]_D²³ +12.0° (c 1, CHCl₃/MeOH = 1:1); mP 124-126°C ; FDMS m/z 397 (M+1)⁺ ; ¹H-NMR (500MHz, CDCl₃/CD₃ = 1:1) δ 4.93-4.96(1H, m, H₂), 3.91(1H, dd, J = 6.7, 12.2Hz), 3.85 (1H, dd, J = 4.9, 12.2Hz), 3.54-3.60(1H, m), 3.50(1H, dd, J = 1.8, 8.5 Hz), 3.19(3H, S), 1.75-1.83(1H, m), 1.53-1.62(1H, m), 1.2-1.45(24H, m), 0.89(3H, t, J = 6.7 Hz).

Synthesis of Compound G7

[0039] To a solution (1000 mL) of Compound G6 (145 g, 365 mmol) in DMF was added sodium azide (47 g, 730 mmol), and the mixture was stirred at 95°C for 4 hours. The reaction solution was concentrated. Ethyl acetate (450 mL) was added to the residue, and the resultant solution was washed with water. The aqueous layer was re-extracted with ethyl acetate several times. All of the organic layers obtained were combined, washed with brine, dried over anhydrous magnesium sulfate and then concentrated under reduced pressure, thereby giving a crude product of Compound G7 in an amount of 122 g (yield: 97%). For the subsequent reaction, the crude product was used without any further purification. For the preparation of a sample for analyses, a portion of the crude product was subjected to further purification by chromatography on a silica gel column with hexane:ethyl acetate (9:1) as an eluent.

[α]_D²³ +16.5° (c 0.5, CHCl₃/MeOH = 1:1); mP 92-93°C ; FDMS m/z 344(M+1)⁺ ; ¹H-NMR (500MHz, CD₃OD) δ 3.91(1H, dd, J = 3.7, 11.6Hz), 3.75(1H, dd, J = 7.9, 11.6 Hz), 3.49-3.61 (3H, m), 1.50-1.71(2H, m), 1.22-1.46(24H,m), 0.90(3H, t, J = 6.7 Hz).

Synthesis of Compound G8

[0040] To a solution (750 mL) of Compound G7 (121 g, about 352 mmol) in methylene chloride were added pyridine (250 mL) and trityl chloride (124 g, 445 mL), and the mixture was stirred at room temperature for 16 hours. Ethanol (30 mL) was added dropwise thereto and then stirred at room temperature for 30 minutes. After washing with a saturated sodium hydrogencarbonate solution, a saturated aqueous ammonium chloride solution and brine, the reaction solution was dried over anhydrous magnesium sulfate and then concentrated under reduced pressure. The residue was purified by chromatography on a silica gel column with hexane:ethyl acetate (10:1) as an eluent, thereby giving a product of Compound G8 in an amount of 34.4 g (yield based on the amount of Compound G6: 52%).

[α]_D²⁴ +11.9° (c 0.9, CHCl₃) ; FDMS m/z 585 M⁺ ; ¹H-NMR (500MHz, CDCl₃ + D₂O) δ 7.24-7.61(15H, m), 3.62-3.66(2H, m), 3.51-3.57(2H, m), 3.42(1H, dd, J = 6.0, 10.4 Hz), 1.23-1.56(26H, m), 0.88(3H, t, J = 6.7 Hz).

Synthesis of Compound G9

[0041] To a solution (300 mL) of Compound G8 (33.5 g, 57.3 mmol) in DMF was added 60% hydrogenated sodium (5.5 g, about 138 mmol in terms of NaH), and the mixture was stirred for 40 minutes. The reaction solution was cooled to 0°C, and then benzyl chloride (15 mL, 120 mmol) was added dropwise thereto. The resultant solution was stirred for

18 hours while gradually elevating the solution temperature to room temperature. Ice-cooled water (100 mL) was added to the reaction solution to stop the reaction, and the reaction solution was then extracted with ethyl acetate. The extract was washed with brine 3 times. All of the organic layers obtained were combined, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure, thereby giving a crude product of Compound G9 in an amount of 42.2 g (yield: 96%). For the subsequent reaction, the crude product was used without any further purification. For the preparation of a sample for analyses, a portion of the crude product was subjected to further purification by chromatography on a silica gel column with hexane:ethyl acetate (100:1) as an eluent.

[α]₂₄^D+9.8° (c 1.0, CHCl₃), FDMS m/z 738 (M-N₂)⁺; ¹H-NMR (500MHz, CDCl₃) δ 7.07-7.48(25H, m), 4.57(1H, d, J = 11.6 Hz), 4.44(1H, d, J = 11.6 Hz), 4.41 (2H, s), 3.73-3.79 (1H, m), 3.46-3.56 (2H, m), 3.37 (1H, dd, J = 8.6, 10.4 Hz), 1.20-1.64 (26H, m), 0.88 (3H, t, J = 6.7 Hz).

Synthesis of Compounds G10 and G11

[0042] To a solution (250 mL) of Compound G9 (41.2 g, about 54 mmol) in 1-propanol was added methanol (30 mL), and subsequently 5% palladium on carbon (4.1 g) and ammonium formate (27.1 g, 4.3 mol) were added thereto. After the mixture was stirred at room temperature for 16 hours, it was diluted with ethyl acetate and then filtered with celite. The filtrate was concentrated under reduced pressure and then dissolved in ethyl acetate. The resultant solution was washed 3 times with a saturated aqueous sodium hydrogencarbonate solution and subsequently with brine. All of the organic layers obtained were combined, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure, thereby giving a crude product of Compound G10 in an amount of 38.9 g (98%). For the subsequent reaction, the crude product (Compound G10) was used without any further purification.

[0043] To a solution (300 mL) of Compound G10 in methylene chloride were added hexacosanic acid (22.4 g, 56.5 mmol) and WSC hydrochloride (12.6 g, 64.6 mmol), and the mixture was refluxed with heating for 2 hours. After cooling to room temperature, the reaction solution was concentrated under reduced pressure. Ethyl acetate (500 mL) was added to the residue, and the resultant solution was washed several times with a 0.5 M aqueous solution of hydrochloric acid, brine, a saturated aqueous sodium hydrogencarbonate solution and then brine. All of the organic layers obtained were combined, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure, thereby giving a crude product of Compound G11 in an amount of 53.2 g (yield: 88%). For the subsequent reaction, the crude product (Compound G11) was used without any further purification. For the preparation of a sample for analyses, a portion of the crude product was subjected to further purification by chromatography on a silica gel column with hexane:ethyl acetate (100:1) as an eluent.

[α]₂₄^D+5.3° (c 0.4, CHCl₃); FDMS m/z 1118 M⁺; ¹H-NMR (500MHz, CDCl₃) δ 7.20-7.38(25H, m), 5.57(1H, d, J = 9.1 Hz), 4.80 (1H, d, J = 11.6 Hz), 4.48-4.50 (3H, m), 4.24-4.32(1H, m), 3.83 (1H, dd, J = 3.0, 6.7 Hz), 3.43-3.51(2H, m, H1a), 3.29(1H, dd, J = 4.3, 9.8 Hz), 1.92(2H, t, J = 7.3 Hz), 1.28-1.60(72H, m), 0.88(6H, t, J = 6.7 Hz).

Synthesis of Compound G12

[0044] To a solution (180 mL) of Compound G11 (52.2 g, about 47 mmol) in methylene chloride was added methanol (36 mL), and a 10% solution of hydrochloric acid in methanol (3.0 mL) was added dropwise thereto. The mixture was stirred at room temperature for 2 hours. The reaction solution was neutralized with sodium hydrogencarbonate powder (18 g) and then filtered with celite. The residue was washed with methylene chloride. The filtrate and the washings were combined, and then washed with brine. The organic layer obtained was dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure. The residue was dissolved in hot acetone and then cooled to 0°C to cause precipitation of a product of Compound G12 in a purified form in an amount of 38.6 g (yield based on the amount of G9: 77%).

[α]₂₄^D-29.7° (c 0.7, CHCl₃); mp 75-76.5°C; FDMS m/z 876M⁺; ¹H-NMR (500MHz, CDCl₃) δ 7.30-7.47(10H, m), 6.03(1H, d, J = 7.9 Hz), 4.72 (1H, d, J = 11.6 Hz), 4.66(1H, d, J = 11.6 Hz), 4.61(1H, d, J = 11.6 Hz), 4.45(1H, d, J = 11.6 Hz), 4.12-4.17(1H, m), 4.00(1H, dt, J₁ = 4.3, J₂ = 7.3 Hz), 3.67-3.72(2H, m), 3.61(1H, ddd, J = 4.3, 8.6, 11.6 Hz), 1.94-2.05(2H, m), 1.15-1.69(72H, m), 0.88(6H, t, J = 6.1 Hz).

Synthesis of Compound G13

[0045]

(1) 2,3,4,6-Tetra-O-benzyl-D-galactopyranosyl acetate (79.8 g) was dissolved in a mixed solution of toluene (160

mL) and isopropyl ether (520 mL) and then cooled to -10 - 0°C. To the solution was added a solution containing 2.0 equivalent amount of HBr in isopropyl ether (2.8 mmol/mL, about 100 mL). After the reaction solution was stirred at -10 - 0°C for about 90 minutes, a 5% aqueous solution of sodium hydrogencarbonate was poured thereto and stirred to neutralize the excess HBr. The whole of the solution was transferred to a separatory funnel to cause separation into an aqueous layer and an organic layer. The aqueous layer obtained was discarded and the organic layer was washed with a 10% aqueous solution of sodium chloride twice. The organic layer was concentrated under reduced pressure, thereby giving a syrup product of 2,3,4,6-tetra-O-benzyl- α -D-galactopyranosyl bromide (Gal-Br). (2) To a solution (420 mL) containing Compound G12 (60.0 g, 68.6 mmol), tetrahexylammonium bromide (89.4 g, 206 mmol) and Molecular Sieves 4A powder (60 g) in toluene were added DMF (140 mL) and subsequently a solution (250 mL) of Gal-Br (about 137 mmol) in toluene. The mixture was stirred at room temperature for 72 hours. Methanol (12 mL) was added to the reaction solution and stirred for 2 hours. After the solution was filtered with celite, the filtrate was washed with a saturated aqueous sodium hydrogencarbonate solution and brine, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure. Acetonitrile was added to the residue, and the resultant solution was stirred for 2 hours to cause precipitation. The precipitate obtained was dried under reduced pressure to give a dry powder product. The powder product was purified by chromatography on a silica gel column with hexane:ethyl acetate (8:1) as an eluent, thereby giving Compound G13 in an amount of 70.9 g (yield: 74%).

$[\alpha]_{24}^{D+18.8^{\circ}}$ (c 0.9, CHCl_3) ; mp 74-75°C ; FDMS m/z 1399 M+1)⁺ ; ¹H-NMR (500MHz, CDCl_3) δ 7.21-7.37(30H, m), 6.12 (1H, d, J = 9.0 Hz), 4.91(1H, d, J = 11.6 Hz), 4.84(1H, d, J = 3.7 Hz), 4.72-4.80 (4H, m), 4.35-4.65 (7H, m), 4.12-4.18 (1H, m), 3.99-4.05(2H, m), 3.84-3.93 (4H, m), 3.73(1H, dd, J = 3.7, 11.0Hz), 3.47-3.51(2H, m), 3.42(1H, dd, J = 6.1, 9.1Hz), 1.87-1.99(2H, m), 1.18-1.70(72H, m), 0.88(6H, t, J = 7.4 Hz).

(2S,3S,4R)-1-O-(α -D-Galactopyranosyl)-N-hexacosanoyl-2-amino-1,3,4-octadecanetriol (KRN7000)

[0046] Compound G13 (60.0 g, 42.9 mmol) was suspended into ethanol (960 mL), and a 20% suspension of palladium hydroxide (6.0 g) in ethanol was added thereto. 4-Methylcyclohexene (120 mL, 93.5 mmol) was added to the suspension as a donor compound for hydrogen, and then refluxed under heating for 4 hours. Thereafter, the resultant solution was filtered to remove the catalyst. The residue was washed with hot ethanol. The filtrate was allowed to stand at room temperature to cause precipitation of a white product. The white precipitate was filtered out and then dried under reduced pressure. The powder product obtained was suspended into ethanol:water (92:8, 3.5 L), dissolved by heating while stirring, and then allowed to stand to cause precipitation again. The precipitate solution was filtered and the filter cake obtained was dried under reduced pressure, thereby giving the title compound in a white powder form in an amount of 35.0 g (yield: 95%).

$[\alpha]_{24}^{D+43.6^{\circ}}$ (c 1.0, pyridine) ; mp 189.5-190.5°C ; negative FABMS m/z 857(M-H)⁻ ; IR(cm^{-1} , KBr) 3300, 2930, 2850, 1640, 1540, 1470, 1070 ; ¹H-NMR(500MHz, $\text{C}_5\text{D}_5\text{N}$) δ 8.47 (1H, d, J = 8.5Hz), 5.58(1H, d, J = 3.7Hz), 5.27(1H, m), 4.63-4.70 (2H, m), 4.56(1H, m), 4.52(1H, t, J = 6.1 Hz), 4.37- 4.47 (4H, m), 4.33(2H, m), 2.45(2H, t, J = 7.3 Hz), 2.25-2.34 (1H, m), 1.87-1.97(2H, m), 1.78-1.85(2H, m), 1.62-1.72(1H, m), 1.26-1.45(66H,m), 0.88(6H, t, J = 6.7Hz). ¹³C-NMR (125MHz, $\text{C}_5\text{D}_5\text{N}$) δ 173.2(s), 101.5(d), 76.7(d), 73.0(d), 72.5(d), 71.6(d), 71.0(d), 70.3(d), 68.7(t), 62.7(t), 51.4(d), 36.8(t), 34.4(t), 32.1(t), 30.4(t), 30.2(t), 30.03(t), 30.00(t), 29.93(t), 29.87(t), 29.81(t), 29.76(t), 29.6(t), 26.5(t), 26.4(t), 22.9(t), 14.3(q).

PHARMACOLOGICAL TESTS

[Pharmacological Test 1] Effect of KRN 7000 on human peripheral blood-derived APC

[0047] This experiment was designed to study an effect of KRN7000 (Compound No.14, a representative glycoside compound of the invention) on APCs prepared from human peripheral blood. The APCs (mainly comprised of dendritic cells) derived from human peripheral blood were prepared essentially according to the method of Hsu et al. (*supra*). Briefly, human peripheral blood was layered over Ficoll-Paque and centrifuged at 2,000 rpm for 35 minutes to give a mononuclear cell fraction. A portion of the mononuclear cell fraction was suspended into PBS (Ca^{++} , Mg^{++} free) supplemented with 5% of inactivated human AB serum. The resultant solution was layered over 50% Percoll, and then centrifuged at 3,000 rpm for 20 minutes to give a high-density fraction. The high-density fraction was panned using a human γ -globulin-coated dish to remove FcR⁺ cells therefrom (i.e., a monocyte-removed fraction). The resultant fraction was reacted with anti-CD3 monoclonal antibodies, anti-CD21 monoclonal antibodies and anti-CD56 monoclonal antibodies. After washing, the resultant solution was panned using an anti-mouse IgG antibody-coated dish to remove cells that were reactive with the above antibodies (i.e., T cells, B cells and NK cells). The cell fraction thus obtained (i.e., a

fraction from which monocytes, T cells, B cells and NK cells were removed) was cultured in RPMI 1640 medium supplemented with 5% of inactivated human AB serum in the presence of a vehicle (DMSO, final concentration: 0.1%) and KRN7000 (final concentration: 100 ng/mL) for 40 hours. After washing, the resultant cells were provided for use as APCs.

5 [0048] On the other hand, as responder cells, CD4⁺ T cells were purified from the rest of the above-prepared mononuclear cell fraction using a human CD4⁺ T cell-enrichment column, and were cultured in RPMI 1640 medium supplemented with 5% of inactivated human AB serum until they were subjected to use with the APCs.

[0049] The APCs (5×10^4 cells/well) and the responder cells (1×10^5 cells/well) both prepared in the above steps were added to wells of a 96-well flat-bottomed plate, and then subjected to autologous MLR assay. During the last 12 hours 10 of the 6-day culture period, tritium-thymidine (³H-TdR) was added to each of the wells (0.5 μ Ci/well), and then the cells were harvested. The amount of the ³H-TdR taken up into the cells was determined using a liquid scintillation counter. The results (mean value and standard deviation of 3 wells) are shown in Fig. 1. In Fig. 1, "V-APC" and "KRN-APC" refer to the APCs pretreated with the vehicle and KRN7000, respectively.

[0050] As shown in Fig. 1, when each of CD4⁺ T cell, the vehicle-pretreated APC (V-APC) and the KRN7000-pretreated APC (KRN-APC) were cultured singly, in either case, the cells took up no or a negligible amount of ³H-TdR. In contrast, when the above-prepared T cell was mixed with V-APC, an obvious ³H-TdR uptake was observed. When the T cell was mixed with KRN-APC, a significantly remarkable ³H-TdR uptake was observed. These results demonstrate that KRN-APC has a more significant autologous MLR enhancing effect than V-APC. Accordingly, KRN7000 has an activating effect on the APCs prepared from human peripheral blood (i.e., an antigen-presenting function enhancing effect). 20

[Pharmacological Test 2] Effect of KRN7000 on human umbilical cord blood-derived APC

[0051] APCs derived from human umbilical cord blood (CD1c⁺ positive cells; CD1c⁺ cells) were prepared in the following manner. That is, human umbilical cord blood was layered over Lymphoprep and centrifuged at 1500 rpm for 20 minutes to give a mononuclear cell fraction. A portion of the mononuclear cell fraction was washed with PBS supplemented with 2% inactivated FBS and 1 mM EDTA twice, and FcR was blocked with human γ -globulin. The solution was reacted with anti-CD1c antibodies, and washed with PBS supplemented with 2% FBS and 1 mM EDTA twice. The solution was further reacted with anti-mouse IgG microbeads, washed with PBS supplemented with 2% FBS and 1 mM EDTA, and then suspended into PBS supplemented with 0.5% BSA and 5 mM EDTA. CD1c⁺ cells were obtained from the suspension using a MiniMACS magnetic column previously washed with PBS supplemented with 0.5% BSA and 5mM EDTA. The CD1c⁺ cells obtained were suspended into RPMI 1640 medium supplemented with 10% inactivated human AB blood serum to adjust its density to 1×10^6 cells/mL. The cell suspension was added to wells of a 24-well plate (2 mL/well), and then a vehicle (DMSO, final concentration: 0.1%) or KRN7000 (final concentration: 100 ng/mL) was added to each of the wells. After incubating for 3 days, the plate was washed 3 times and provided for use as APCs. 35

[0052] CD4⁺ T cells were purified from the rest of the above-prepared mononuclear cell fraction using a human CD4⁺ T cell-enrichment column. The T cells obtained were suspended into RPMI 1640 medium supplemented with 10% inactivated human AB blood serum to adjust its density to 1×10^6 cells/well. The suspension was added to wells of a 24-well plate (2 mL/well) and cultured for 3 days. The T cells thus obtained were used as responder cells for autologous MLR assay. On the other hand, a peripheral blood was collected from a normal volunteer, and CD4⁺ T cells were purified therefrom using a human CD4⁺ T cell-enrichment column. The T cells thus obtained was used as responder cells for allogeneic MLR assay. The responder cells (1×10^5 cells/50 mL/well) and the stimulator cells (0, 1000, 10000 cells/50 mL/well) were added to wells of a 96-well flat-bottomed plate, and the plate was incubated. For allogeneic MLR assay, the plate was incubated for 4 days, ³H-TdR was added to each of the wells (0.5 μ Ci/well). After 20 hours, the cells were harvested and the amount of the ³H-TdR taken up into the cells was determined using a liquid scintillation counter. The results (mean value and standard deviation of 3 wells) are shown in Fig. 2. 45

[0053] For autologous MLR assay, the plate was incubated for 7 days, ³H-TdR was added to each of the wells (0.5 μ Ci/well). After 8 hours, the cells were harvested and the amount of the ³H-TdR taken up into the cells was determined using a liquid scintillation counter. The results (mean value and standard deviation of 3 wells) are shown in Fig. 3. In Figs. 2 and 3, "V-APC" and "KRN-APC" refer to the APCs (CD1c⁺ cells) pretreated with the vehicle and KRN7000, respectively. 50

[0054] As shown in Fig. 2., the CD1c⁺ cells stimulated with the vehicle (APCs; V-APC) promoted the uptake of ³H-TdR (proliferation of allogeneic T cells) in a manner dependent on the number of the V-APC, whereas the CD1c⁺ cells stimulated with KRN7000 (APCs; KRN-APC) showed a stronger allogeneic T cell-proliferation-promoting effect than V-APC.

55 [0055] As shown in Fig. 3, when autologous T cells were used as the responder cells, the CD1c⁺ cells stimulated with the vehicle (V-APC) hardly stimulated the proliferation of the autologous T cells even if the APCs were added to the wells in an amount of 10000 cells/well; whereas the CD1c⁺ cells treated with KRN7000 (KRN-APC) significantly stimulated the proliferation of the autologous T cells in a manner dependent on the number of the APCs, similar to the case

for the allogeneic T cells.

[Pharmacological Test 3] Effect of α - and β -galactosylceramides and α - and β -glucosylceramides on murine spleen-derived APC

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[0056] The preparation of dendritic cell-rich APCs from a murine spleen was performed essentially in accordance with the method of Clowley, M. et al. (*supra*). Briefly, the spleen was removed from a BDF1 mouse. After the treatment with 100 U/mL of collagenase, the spleen was dissected with tweezers to separate into a cell suspension and a tissue debris. The tissue debris was suspended into a 400 U/mL solution of collagenase. After culturing in a CO₂ incubator for 20 minutes, the suspension was passed through a stainless steel mesh with an inner syringe to give a cell suspension. Two of the cell suspensions were combined, and subjected to centrifugation. The cell suspension was subjected to density gradient centrifugation with a high-density BSA to give a low-density cell fraction. The cell fraction was plated onto a 60 mm dish and cultured for 2 hours. The procedure for removing the floating cells was repeated 3 times. RPMI 1640 medium supplemented with 10% inactivated FBS was added to the dish and subsequently any one of KRN7000 (Compound No.14), AGL583 (a β -anomer of KRN7000), AGL517 (Compound No.3), AGL564 (a β -anomer of AGL517), AGL563 (Compound No.4), AGL562 (a β -anomer of AGL-563) and a vehicle (DMSO, final concentration: 0.1%) was also added thereto to a final concentration of 100 ng/mL. After culturing overnight, the non-adherent cells were collected, and the resultant cells were provided for use as APCs.

[0057] On the other hand, BDF1 mouse spleen cells were treated with a red blood cell-lysing buffer comprised of NH₄Cl and Tris-HCl to hemolyze red blood cells, and the resultant spleen cells were suspended into RPMI 1640 medium supplemented with 10% inactivated FBS. The spleen cells were inoculated onto a 100 mm dish, and cultured in a CO₂ incubator for 2 hours. Thereafter, the non-adherent cells were recovered, which were provided for use as responder cells.

[0058] The above-prepared APCs (1×10^3 , 3.3×10^3 or 1×10^4 cells/well) and the responder cells (2.5×10^5 cells/well) were added to wells of a 96-well plate for syngeneic MLR assay. After culturing for 1 day, ³H-TdR was added to each of the wells (0.5 μ Ci/well). After 16 hours, the cells were harvested, and the amount of ³H-TdR taken up into the cells was determined using a liquid scintillation counter. The results (mean value and standard deviation of 3 wells) are shown in Fig. 4.

[0059] In Fig. 4, "V-APC", "KRN-APC", "583-APC", "517-APC", "564-APC", "563-APC" and "562-APC" refer to the APCs pretreated with the vehicle, KRN7000, AGL-583, AGL-517, AGL-564, AGL-563 and AGL-562, respectively.

[0060] As shown in Fig. 4, the APCs pretreated with α -galactosylceramides such as KRN-APC and 517-APV showed a significant syngeneic MLR enhancing effect, whereas the APCs pretreated with β -galactosylceramides such as 583-APC and 564-APC did not show any such effect. The APCs pretreated with α -glucosylceramides such as 563-APC also showed a remarkable syngeneic MLR enhancing effect, whereas the APCs pretreated with β -glucosylceramides such as 562-APC did not show any such effect.

[Pharmacological Test 4] APC therapy with murine spleen-derived APCs pretreated with KRN7000

[0061] To study an anti-tumor effect of murine spleen-derived APCs activated with KRN7000 on mice implanted with tumor cells, experiments were performed with BDF1 mice (6-week-old, female). All experiments included 6 mice per group. Murine T cell lymphoma EL-4 cells were injected intravenously into each mouse at a level of 1×10^5 cells/mouse (injection day: day 0). On day 1, the vehicle-pretreated APCs (V-APC) and the KRN7000-pretreated (100 ng/mL) APCs (KRN-APC) both prepared in the same manner as in Pharmacological Test 3 were each injected intravenously into the mice at a level of 5×10^5 cells/mouse. As a positive control, KRN7000 was injected intravenously into the mice at a dose of 100 μ g/kg on days 1, 5 and 9. Each mouse was checked for a sign of life daily to determine its survival period. The results are shown in Fig. 5.

[0062] As shown in Fig. 5, when V-APC was injected, no prolongation of survival period was observed. However, when KRN-APC was injected, a remarkable increase in survival period was observed and 50% of the KRN-APC-injected mice were cured completely. When KRN7000 was injected intravenously at a dose of 100 μ g/kg 3 times, a remarkable increase in survival period was observed, however, it was found that the potency of the effect was weaker than that when KRN-APC was injected only once.

[Pharmacological Test 5] Antigen presenting function enhancing effect of KRN7000 on murine bone marrow-derived dendritic cell

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[0063] Dendritic cell-rich APCs derived from murine bone marrow were prepared according to the method of Inaba, K. et al. (*supra*) with some modifications (Yamaguchi Y. et al., Stem Cells, 1977:15:144-153). Briefly, bone marrow cells from BALB/c mice were prepared. After hemolyzing the red blood cells with a NH₄Cl solution, FcR⁺ cells were removed

by panning using a human γ -globulin-coated dish. The cells thus obtained were suspended in RPMI medium supplemented with 10% FCS, and cultured in the presence of 10 ng/mL of mouse rGM-CSF and 10 ng/mL of human rTGF- β for 6 days at 5×10^5 cells/well (1 ml/well, a 24-well plate). Every 2 days, each well was washed briefly using a Pasteur pipette and then about 75% of the medium was sucked out and supplemented with 1 ml of a fresh medium containing the above factors. After culturing for 6 days, non-adherent cells were recovered, and FcR $^+$ cells were removed therefrom by panning using a human γ -globulin-coated dish. The cells thus prepared were cultured in a culture medium supplemented with 10 ng/mL of mouse rGM-CSF and 10 ng/mL of human rTNF- α for additional 2 days. During this culture, either a vehicle (DMSO, final concentration: 0.1%) or KRN7000 (final concentration: 100 ng/mL) was added to the culture medium. The cells were collected and washed 3 times to provide for use as APCs. On the other hand, as responder cells, T cells were prepared from a BALB/c murine spleen using a T cell enrichment column (R&D). T cells were added to wells of a 96-well plate (3×10^5 cells/well), and the above-prepared APCs were further added thereto for syngeneic MLR assay (3×10^4 , 1×10^4 , 3×10^3 and 1×10^3 cells/well, respectively). After culturing for 2 days, ^3H -TdR was added to each of the wells (0.5 $\mu\text{Ci/mL}$). After 6 hours, the cells were harvested and determined on the uptake of ^3H -TdR into the cells using a liquid scintillation counter. The results (mean value and standard deviation of 3 wells) are shown in Fig. 6. In Fig. 6, "V-APC" and "KRN-APC" refer to the APCs pretreated with the vehicle and KRN7000, respectively.

[0064] As shown in Fig. 6, the murine bone marrow-derived APCs which were stimulated with KRN7000 (KRN-APC) showed a remarkable syngeneic MLR enhancing effect compared to the APCs stimulated with the vehicle (V-APC).

[Pharmacological Test 6] APC therapy with murine bone marrow derived-APCs pretreated with KRN7000

[0065] The experiment was performed to study an anti-tumor effect of murine bone marrow-derived APCs activated with KRN7000 on mice implanted with tumor cells. All experiments were performed with CDF1 mice (6-week-old, female), which were divided into groups consisting of 5 mice. Murine colon adenocarcinoma Colon26 cells were implanted into each mouse intrasplenically at a level of 2×10^6 cells/mouse (inoculation day: day 0). On day 1, the vehicle-pretreated APCs (V-APC) and the KRN7000-pretreated (100 ng/mL) APCs (KRN-APC) both prepared in the same manner as in Pharmacological Test 5 were individually injected intravenously to the mice at a level of 8×10^5 cells/mouse. As a positive control, KRN7000 was injected intravenously to the mice at a dose of 100 $\mu\text{g/kg}$ on days 1, 5 and 9. On day 14, the liver was removed from each of the mice and weighed. The results (mean value and standard deviation of the weights of the liver of 5 mice) are shown in Fig. 7. Here, the liver from the mouse that had not been implanted with the tumor cells was about 1 g. Accordingly, the substantial tumor weight was determined by subtracting 1 g from the weight of the liver of the tumor-implanted mouse.

[0066] As shown in Fig. 7, when V-APC was injected, a slight suppressive effect on tumor growth was observed; whereas when KRN-APC was injected, a remarkable inhibitory effect on tumor growth was observed and no tumor nodule in the liver was observed by naked eyes in 3 of the 5 mice. When KRN7000 was injected intravenously 3 times at a dose of 100 $\mu\text{g/kg}$, a remarkable inhibitory effect on tumor growth was observed, which was at the same level as that when KRN-APC was injected once.

[Pharmacological Test 7] Antigen presenting function enhancing effect of KRN7000 on murine epidermis-derived APCs

[0067] Langerhans's cell-rich APCs were prepared from murine auricle essentially in accordance with the method of Witmer-Pack, M. et al. (*supra*). Briefly, the epidermis of auricle of a BALB/c mouse was peeled into front and back epidermal sheets, and each of the epidermal sheets was soaked into Hanks's solution supplemented with 1% trypsin at 37°C for 30 minutes to 1 hour. The epidermal sheets obtained were placed on a mesh, and shaken up and down in Hanks's solution supplemented with 10% FCS. The cells detached from the epidermal sheets were collected, and then suspended into RPMI solution supplemented with 10% FCS at 10^6 cells/mL. After addition of either a vehicle (DMSO, final concentration: 0.1%) or KRN7000 (final concentration: 100 ng/mL), the cells were cultured under 5% CO_2 at 37°C for 3 days. Thereafter, the cells that were non-adherent to the dish were collected and layered over Lympholite-M. After centrifugation at 1400 rpm for 10 minutes, the low-density cells were collected, washed with RPMI twice, which were provided for use as APCs.

[0068] As responder cells, spleen cells were prepared from BALB/c mice in the same manner as described in Pharmacological Test 3. The responder cells thus obtained were added to wells of a 96-well plate (2.5×10^5 cells/well), and the APCs were also added thereto for syngeneic MLR assay (4×10^4 , $4/3 \times 10^4$ and $4/9 \times 10^4$ cells/well, respectively). After culturing for 2 days, ^3H -TdR was added to each well (0.5 $\mu\text{Ci/mL}$). After 6 hours, the cells were harvested and determined on the amount of ^3H -TdR taken up into the cells using a liquid scintillation counter. The results (mean value and standard deviation of 3 wells) are shown in Fig. 8. In Fig. 8, "V-APC" and "KRN-APC" refer to the APCs pretreated with the vehicle and KRN7000, respectively.

[0069] As shown in Fig. 8, the murine epidermis-derived APCs stimulated with KRN (KRN-APCs) showed a more remarkable syngeneic MLR enhancing effect compared to those stimulated with the vehicle (V-APCs).

[Pharmacological Test 8] APC therapy with murine spleen-derived APCs pretreated with KRN7000 and tumor antigen

[0070] The experiment was performed with BDF1 mice (6-week-old, female), which were divided into groups consisting of 6 mice. Murine melanoma B16 cells were implanted subcutaneously into each mouse at a level of 1.5×10^6 cells/mouse (implantation day: day 0). Several types of pretreated APCs were prepared in the same manner as in Pharmacological Test 3, in which APCs were pretreated with a vehicle (DMSO, final concentration: 0.1%) (V-APC), with both the vehicle (DMSO, final concentration: 0.1%) and a B16-tumor cell lysate (V-T-APC), with KRN7000 (final concentration: 100 ng/mL) (KRN-APC), and with both KRN7000 (final concentration: 100 ng/mL) and a B16-tumor cell lysate (KRN-T-APC), respectively. Each of the pretreated APCs was injected intravenously into the mice at a dose of 5×10^5 cells/mouse on day 1. As a positive control, KRN7000 was injected intravenously into the mice at a dose of 100 μ g/kg on days 1, 5 and 9. The volume of the subcutaneous tumor in each mouse was measured. The results (the average of the tumor volumes of 6 mice) are shown in Fig. 9.

[0071] As shown in Fig. 9, little or no tumor growth-suppressive effect was observed when V-APCs and V-T-APCs were injected. On the contrary, when KRN-APCs were injected once, a tumor growth-inhibitory effect was observed, which was at the same level as that when KRN7000 was injected 3 times intravenously at a dose of 100 μ g/kg. It is interesting to note that the one injection of KRN-T-APC produced a more remarkable tumor growth inhibitory effect than the one injection of KRN-APC or the three intravenous injections of KRN7000 at a dose of 100 μ g/kg.

[Pharmacological Test 9] Effect of KRN7000 on human peripheral blood-derived APCs

[0072] A monocyte fraction was prepared from human peripheral blood mononuclear cells by Percoll density-gradient centrifugation. The monocyte fraction was cultured in the presence or absence of GM-CSF (50 ng/mL) and IL-4 (100 ng/mL), and simultaneously either a vehicle (v; DMSO, 0.1%) or KRN7000 (KRN; 100 ng/mL) was added thereto. After culturing for 3 days, the cells were collected and washed with the medium 3 times to give V-APC, KRN-APC, V-APC-GM and KRN-APC-GM (where "GM" refers to the APCs cultured in the presence of GM-CSF and IL-4), which were then provided for use as APCs. As responder cells, human peripheral blood-derived FcR γ cells were used to perform allogeneic MLR assay.

[0073] As shown in Fig. 10, all of the APCs enhanced the allogeneic MLR response in a manner dependent on the number of the APCs, however, KRN-APC and KRN-APC-GM showed a clearly stronger stimulatory effect on the proliferation of responder cells compared to V-APC and V-APC-GM, respectively. Furthermore, it was also found that the effect of KRN-APC-GM was stronger than KRN-APC. It is assumed that the suppressed proliferation of the responder cells by the addition of KRN-APC-GM in an amount of 20000 cells/well is due to the over growth of the responder cells.

[Pharmacological Test 10] Effect of KRN7000 on human peripheral blood-derived APC

[0074] To study the effect of long-term culture of APC, the following experiment was performed. A monocyte fraction was prepared from human peripheral mononuclear cells by Percoll density-gradient centrifugation. The monocyte fraction was cultured in the presence of GM-CSF (50 ng/mL) and IL-4 (100 ng/mL), and simultaneously either a vehicle (V; 0.1% DMSO) or KRN7000 (KRN; 100 ng/mL) was added thereto. After culturing for 7 days, the cells were recovered and washed with the medium 3 times to give V-APC-GM and KRN-APC-GM, respectively, which were provided for use as APCs. As responder cells, human peripheral blood-derived FcR γ cells were used to perform autologous MLR assay. [0075] As shown in Fig. 11, V-APC-GM did not stimulate the proliferation of the responder cells, whereas KRN-APC-GM enhanced the autologous MLR response in a manner dependent on the number of the APCs.

[Pharmacological Test 11] Effect of KRN7000 on human peripheral blood-derived APC

[0076] To study the effect of long-term culture of APC, the following experiment was performed. A monocyte fraction was prepared from human peripheral mononuclear cells by Percoll density-gradient centrifugation. The monocyte fraction was cultured in the presence of GM-CSF (50 ng/mL) and IL-4 (100 ng/mL) with or without monocyte conditioned medium (MCM) prepared by the method of Bender et al (Bender A. et al., 1996, J. Immunol. Method., 196, 121). Simultaneously either a vehicle (V; 0.1% DMSO) or KRN7000 (KRN; 100 ng/mL) was added to the fraction. After culturing for 11 days, the cells were recovered and washed with the medium 3 times to give V-APC-GM (MCM-), KRN-APC-GM (MCM-), V-APC-GM (MCM+) and KRN-APC-GM (MCM+), respectively, which were provided for use as APCs. As responder cells, human peripheral blood-derived FcR γ cells were used to perform autologous MLR assay.

[0077] As shown in Fig. 12, V-APC-GM (MCM-) did not stimulate the proliferation of the responder cells, whereas KRN-APC-GM (MCM-) enhanced the autologous MLR response in a manner dependent on the number of the APCs. Both V-APC-GM (MCM+) and KRN-APC-GM (MCM+) enhanced the autologous MLR response in a manner dependent on the number of the APCs. The effect of KRN-APC-GM (MCM+) was stronger than that of V-APC-GM (MCM+).

[0078] From the results of Pharmacological Tests 9, 10 and 11 above, it has been found that KRN7000 can activate the APCs from human peripheral blood by various methods.

[Pharmacological Test 12] Effect of KRN7000 on human umbilical cord blood-derived APC

[0079] A monocyte fraction was prepared from human umbilical cord blood mononuclear cells by Percoll density-gradient centrifugation. The monocyte fraction was cultured in the presence of GM-CSF (50 ng/mL) and IL-4 (100 ng/mL), and simultaneously either a vehicle (V; 0.1% DMSO) or KRN7000 (KRN; 100 ng/mL) was added thereto. After culturing for 3 days, the cells were recovered and washed with the medium 3 times to give V-APC-GM and KRN-APC-GM, respectively, which were provided for use as APCs. As response cells, human umbilical cord blood-derived FcR γ cells were used for autologous MLR assay.

[0080] As shown in Fig. 13, V-APC-GM did not stimulate the proliferation of the responder cells, whereas KRN-APC-GM enhanced the autologous MLR response in a manner dependent on the number of the APCs. It is assumed that the suppressed proliferation of the responder cells by the addition of KRN-APC-GM in an amount of 20,000 cells/well is due to the over growth of the responder cells.

[0081] This result shows that KRN7000 can activate the APCs derived from human umbilical cord blood by various methods.

[Pharmacological Test 13] Effect of α -galactosylceramides on murine spleen-derived APC

[0082] To demonstrate that the α -galactosylceramide derivatives of the present invention other than KRN7000 also have an APC-activating effect, the following pharmacological test was performed.

[0083] The dendritic cell-rich APCs were prepared from murine spleen in the same manner as described in Pharmacological Test 3. Briefly, the APCs were treated with any one of KRN7000 (Compound No. 14 of the invention), AGL512 (Compound No. 10 of the invention), AGL525 (Compound No. 16 of the invention), AGL506 (Compound No. 1 of the invention), AGL514 (Compound No. 2 of the invention), AGL571 (Compound No. 5 of the invention) and a vehicle (DMSO, final concentration: 0.1%) at the final concentration of 100 ng/mL. Each of the APCs thus treated (1×10^4 cells/well) and the responder cells (2.5×10^5 cells/well) were added to wells of a 96-well plate and then subjected to syngeneic MLR assay. After culturing for 2 days, ^3H -TdR was added to each well ($0.5 \mu\text{Ci/well}$). After 8 hours, the cells were harvested, and the amount of the ^3H -TdR taken up into the cells was determined using a liquid scintillation counter. The results are shown in Fig. 14.

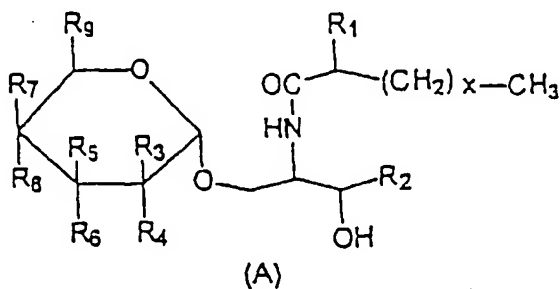
[0084] In Fig. 14, "V-APC", "KRN-APC", "512-APC", "525-APC", "506-APC", "514-APC" and "571-APV" refer to the APCs pretreated with the vehicle, KRN7000, AGL-512, AGL-525, AGL-506, AGL-514 and AGL-571, respectively.

[0085] As shown in Fig. 14, the APCs pretreated with α -galactosylceramide derivatives showed a remarkable syngeneic MLR enhancing effect.

[0086] This result shows that the compounds of the present invention (α -galactosylceramides and α -glucosylceramides) other than KRN7000 also have an effect of activating the dendritic cell-rich APCs derived from murine spleen (i.e., an APC function-enhancing effect).

40 Claims

1. A method for activating human antigen-presenting cells which comprises culturing human-derived antigen-presenting cells *in vitro* with at least one of the glycoside compounds represented by formula (A) or salts thereof:



wherein:

R₁ is H or OH;

X is an integer of from 7 to 25;

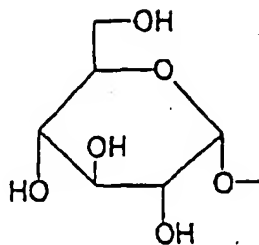
R₂ is a substituent defined by any one of the following (a) to (e):

- (a) -CH₂(CH₂)_YCH₃;
- (b) -CH(OH)(CH₂)_YCH₃;
- (c) -CH(OH)(CH₂)_YCH(CH₃)₂;
- (d) -CH=CH(CH₂)_YCH₃; and
- (e) -CH(OH)(CH₂)_YCH(CH₃)CH₂CH₃;

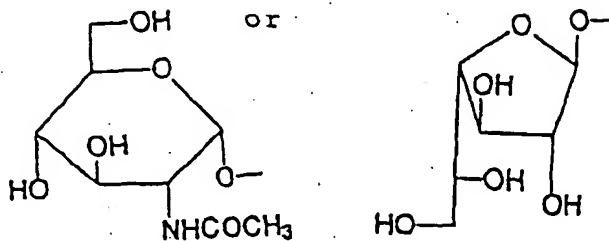
wherein Y is an integer of from 5 to 17;

R₃ is H;

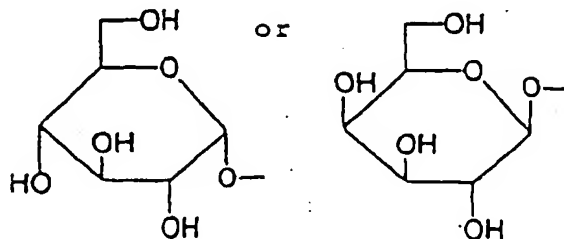
R₄ is OH, NH₂, NHCOCH₃ or



one of R₅ and R₆ is H and the other is OH,

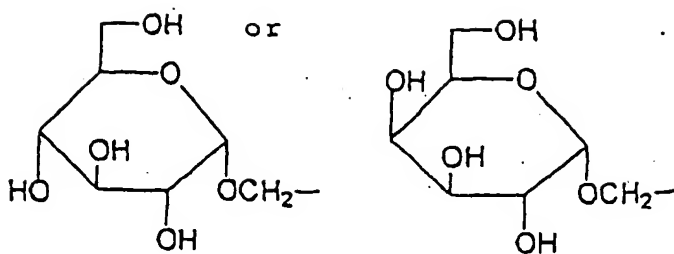


one of R₇ and R₈ is H and the other is OH,

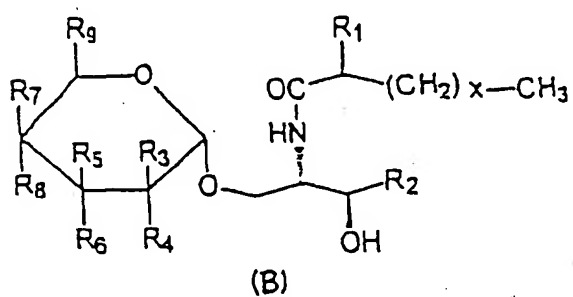


and

R₉ is H, CH₃, CH₂OH,



15 2. The method of claim 1, wherein the glycoside compound is a compound represented by formula (B):

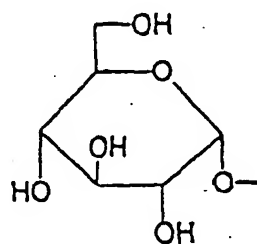


30 wherein:

R_1 , x and R_2 are as defined as in claim 1; and
 R_3 to R_9 are substituents defined by any one of the following (i) to (iii):

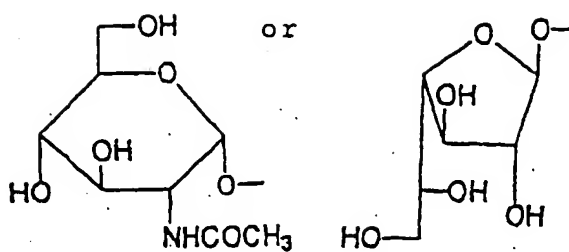
35 (i) each of R_3 , R_6 and R_8 is H;

R_4 is OH or

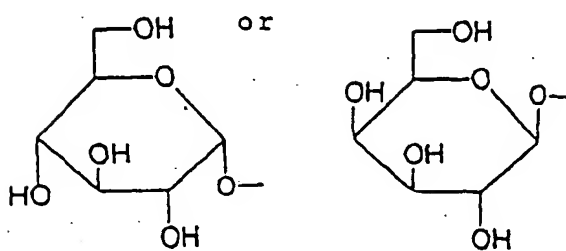


50 R_5 is OH,

55

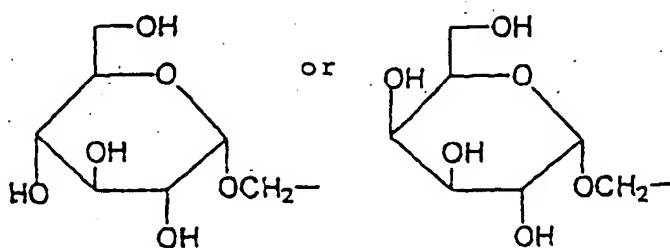


15 R_7 is OH,



and

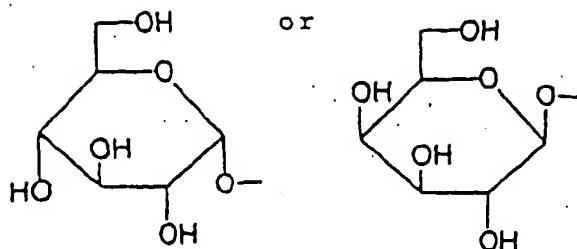
30 R_9 is H, CH_3 , CH_2OH ,



45 (ii) each of R_3 , R_6 and R_7 is H;

R_4 , R_5 and R_9 are as defined as in (i); and

R_8 is OH,



or

15 (iii) each of R_3 , R_5 and R_7 is H;

each of R_4 , R_6 and R_8 is OH; and
 R_9 is H, CH_3 or CH_2OH .

- 20 3. The method of claim 2, wherein, in the glycoside compound, each of R_3 , R_6 and R_8 is H, each of R_4 , R_6 and R_7 is OH, and R_9 is CH_2OH .
4. The method of claim 2 or 3, wherein, in the glycoside compound, R_2 is any one of the substituents (b), (c) and (e).
- 25 5. The method of claim 4, wherein, in the glycoside compound, R_1 is H and R_2 is the substituent (b).
6. The method of claim 5, wherein, in the glycoside compound, the OH group in the substituent (b) is of R configuration.
- 30 7. The method of claim 5 or 6, wherein, in the glycoside compound, X is an integer of 21 to 25 and Y is an integer of 11 to 15.
8. The method of claim 6 or 7, wherein the glycoside compound is (2S,3S,4R)-1-(α -D-galactopyranosyloxy)-2-hexacosanoylamino-3,4-octadecanediol.
- 35 9. The method of any one of claims 1 to 8, wherein the concentration of the glycoside compound or the salt thereof used in the culture is within the range from 0.1 ng/mL to 10 $\mu\text{g/mL}$.
10. The method of claim 9, wherein the concentration of the glycoside compound or the salt thereof is within the range from 0.01 $\mu\text{g/mL}$ to 1 $\mu\text{g/mL}$.
- 40 11. The method of any one of claims 1 to 10, wherein the culture is performed in the presence of GM-CSF and IL-4.
12. The method of claim 11, wherein the culture is performed in the presence of the monocyte conditioned medium (MCM).
- 45 13. The method of any one of claims 1 to 12, wherein the culture is performed in the presence of a tumor antigen.
14. An activated human antigen-presenting cell prepared by activating human-derived antigen-presenting cells by the method of any one of claims 1 to 13.
- 50 15. The activated human antigen-presenting cell of claim 14, wherein the human-derived antigen-presenting cell is selected from the group consisting of a CD1c positive cell, a human monocyte and a human dendritic cell.
- 55 16. The activated human antigen-presenting cell of claim 15, wherein the human-derived antigen-presenting cell is prepared from human peripheral blood.
17. The activated human antigen-presenting cell of claim 15, wherein the human-derived antigen-presenting cell is pre-

pared from human umbilical cord blood.

- 5
18. The activated human antigen-presenting cell of claim 15, wherein the human-derived antigen-presenting cell is prepared from a human bone marrow cell.
19. The activated human antigen-presenting cell of claim 15, wherein the human-derived antigen-presenting cell is prepared from a human epidermis.
- 10
20. An agent for *in vitro* activation of human antigen-presenting cell which comprises, as an active ingredient, at least one of the glycoside compounds or salts thereof of claim 1.
21. The agent of claim 20, wherein at least one of the glycoside compounds or salts thereof of claim 1 is used in combination with a tumor antigen.
- 15
22. A method for treating cancer and infectious diseases including AIDS using the activated human antigen-presenting cell of any one of claims 14 to 19.
- 20
23. A use of the human activated human antigen-presenting cell of any one of claims 14 to 19 in the preparation of a medicine for treating cancer and infectious diseases including AIDS by application of an antigen-presenting cell therapy.
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- 55

Fig. 1

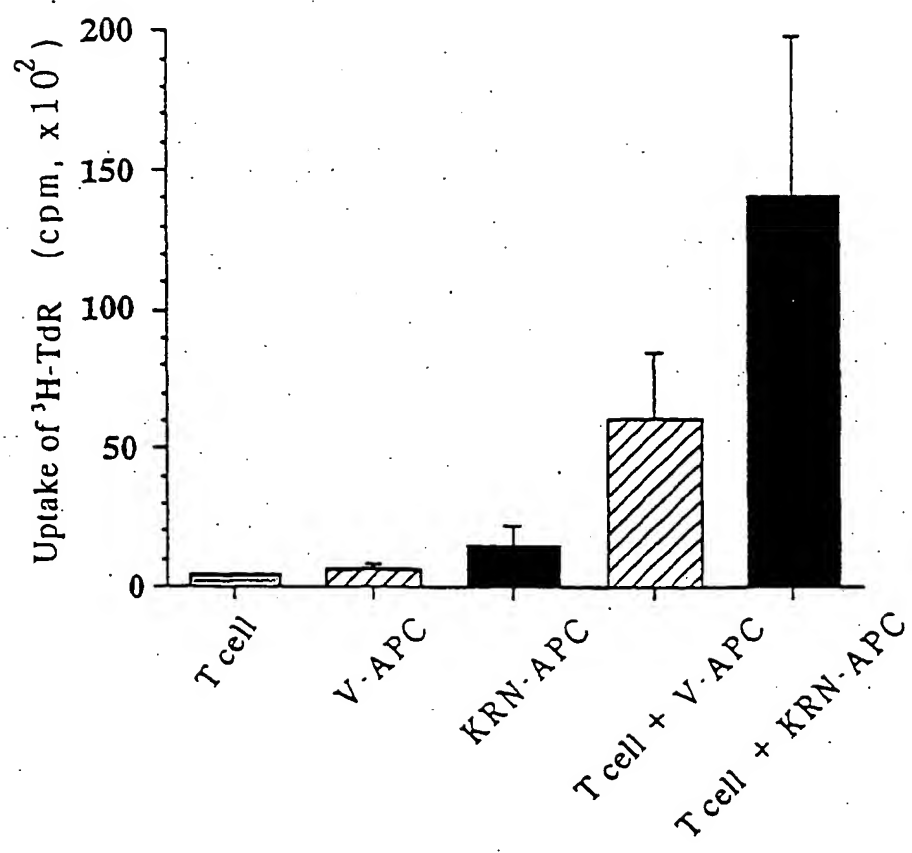


Fig. 2

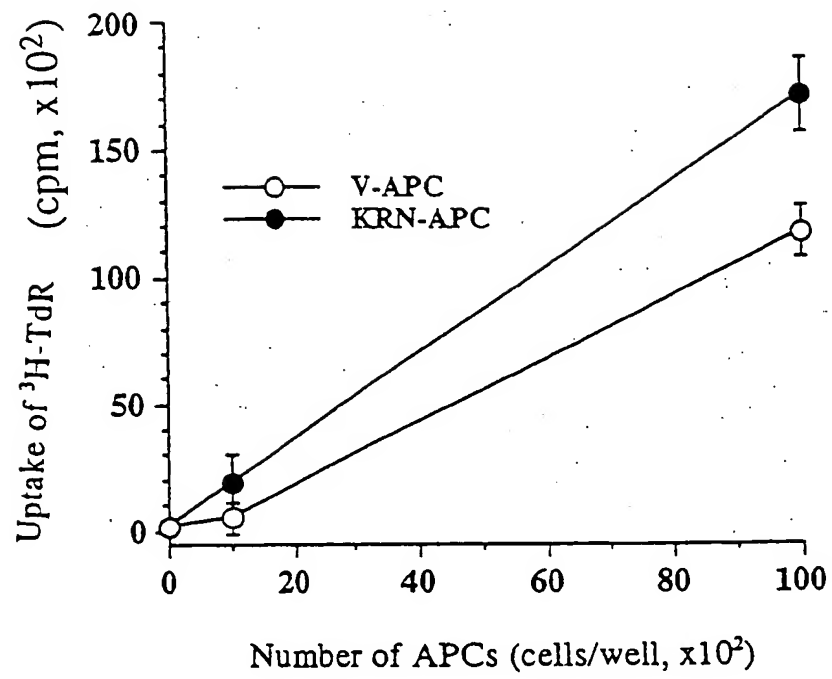


Fig. 3

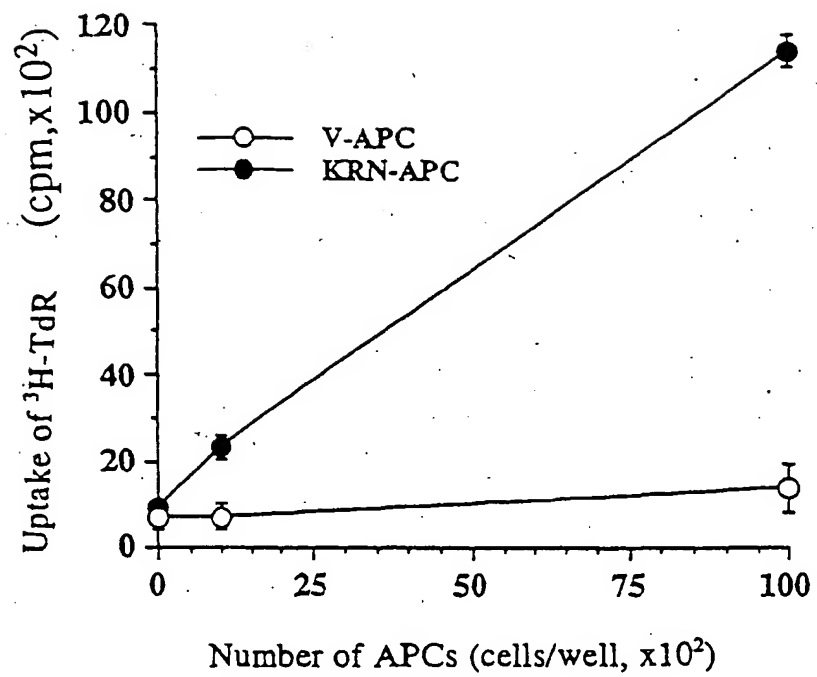


Fig. 4

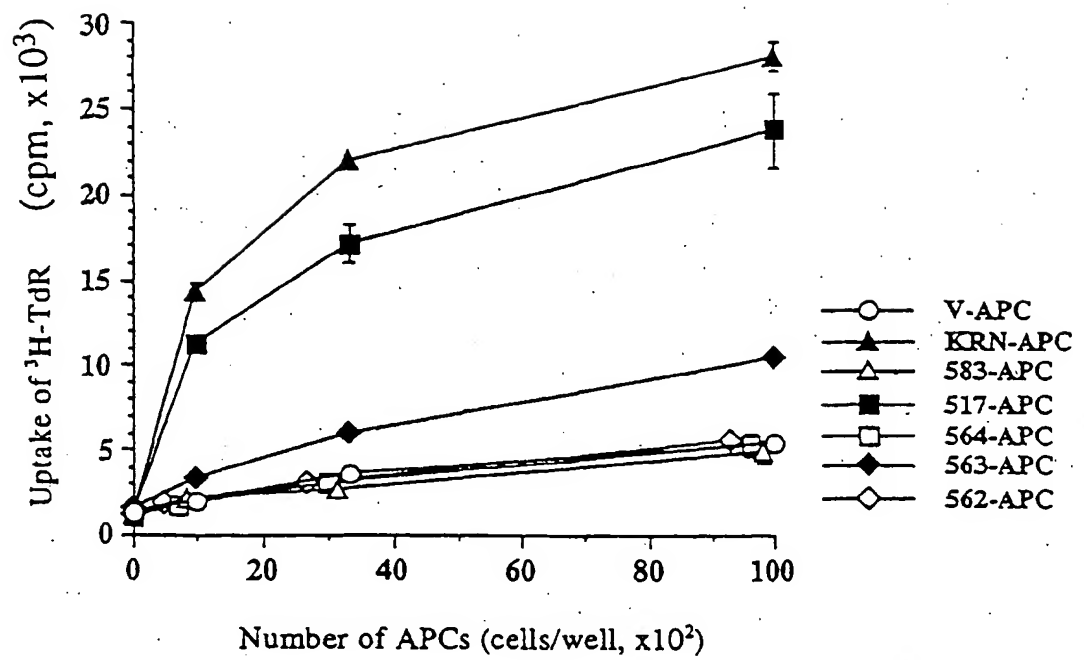


Fig. 5

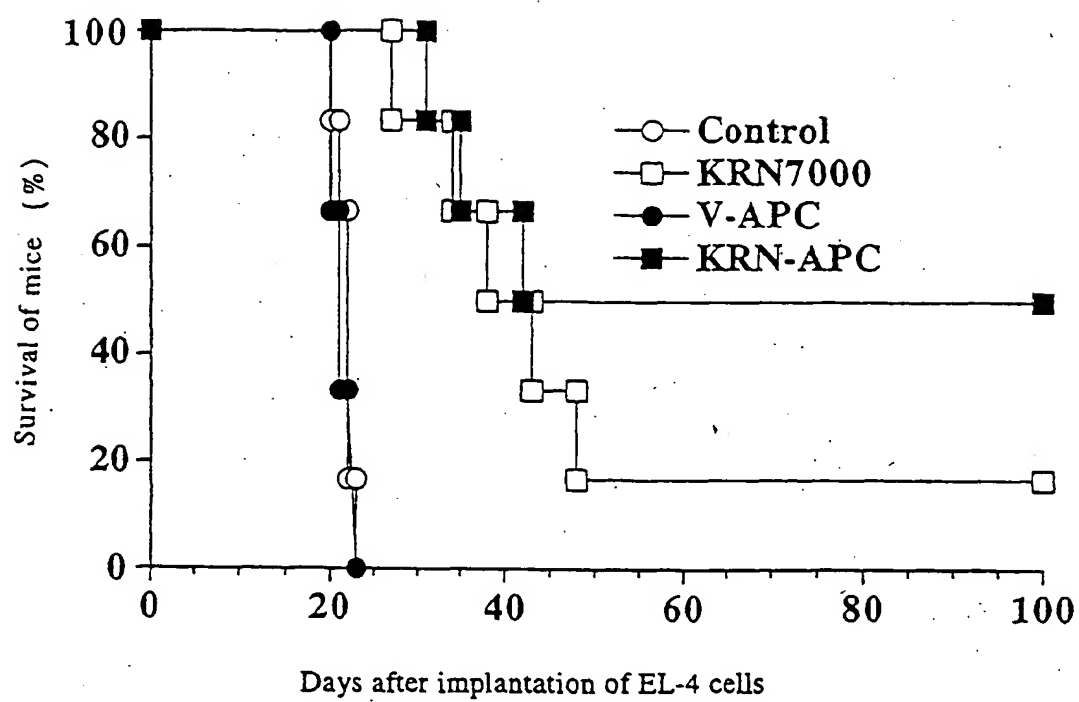


Fig. 6

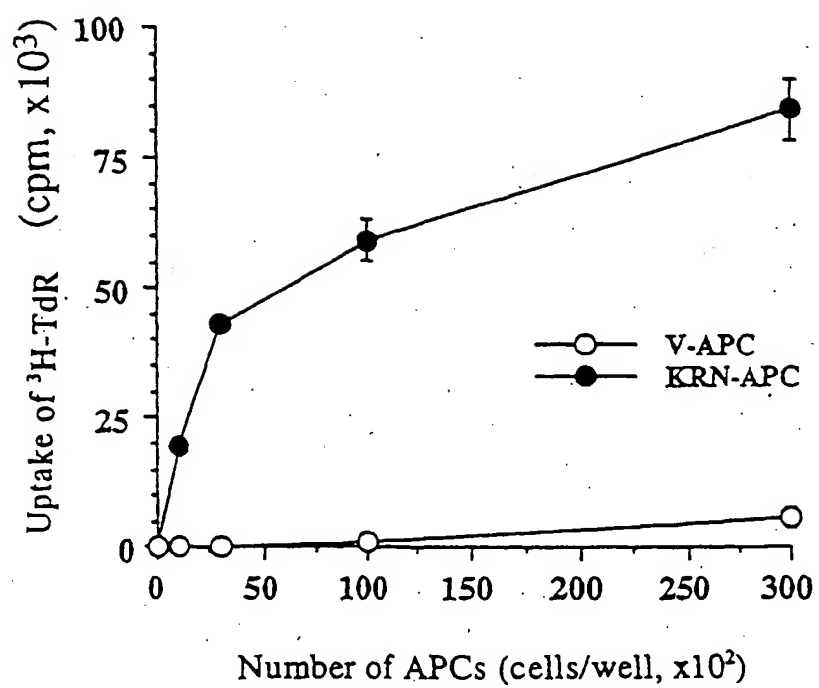


Fig. 7

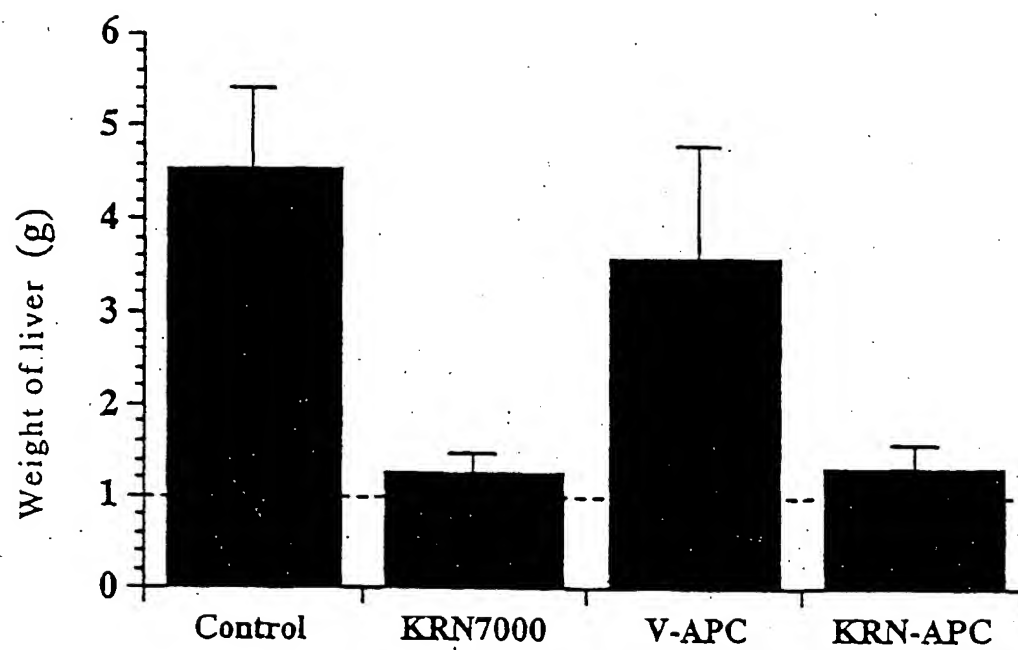


Fig. 8

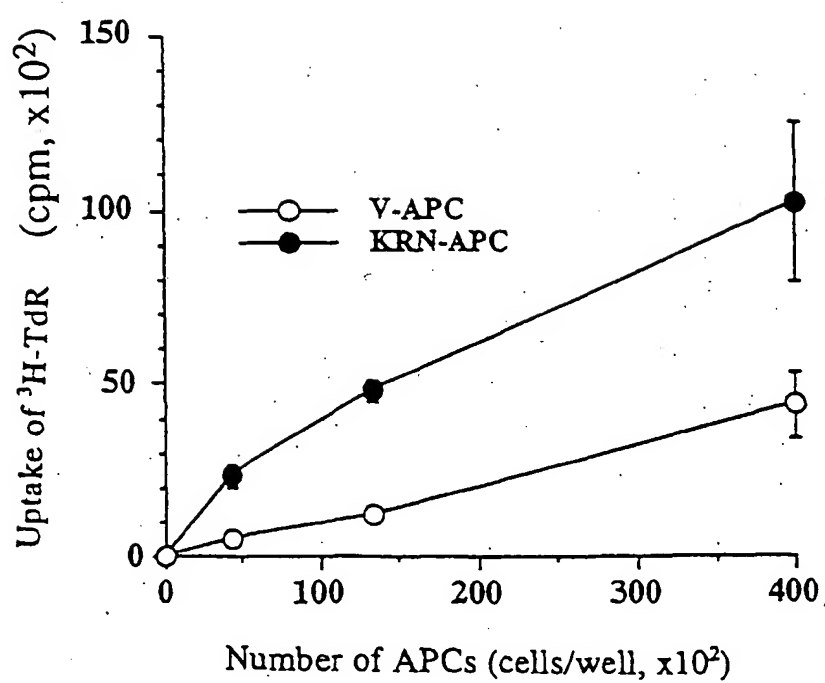


Fig. 9

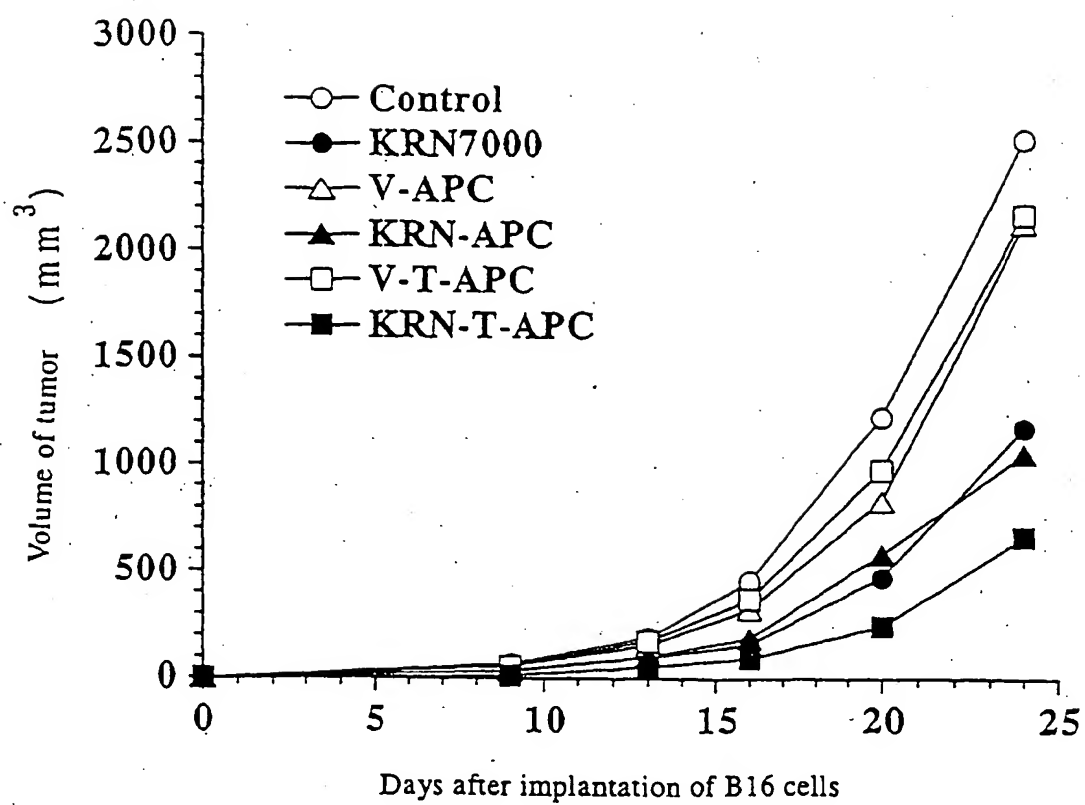


Fig. 10

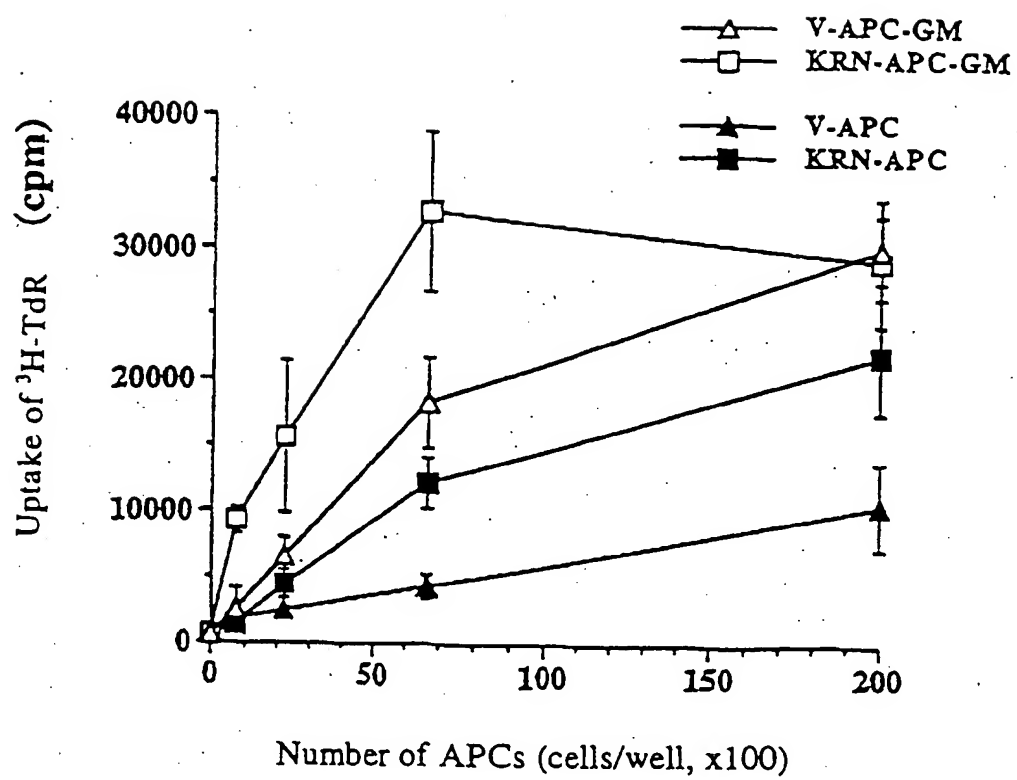


Fig. 11

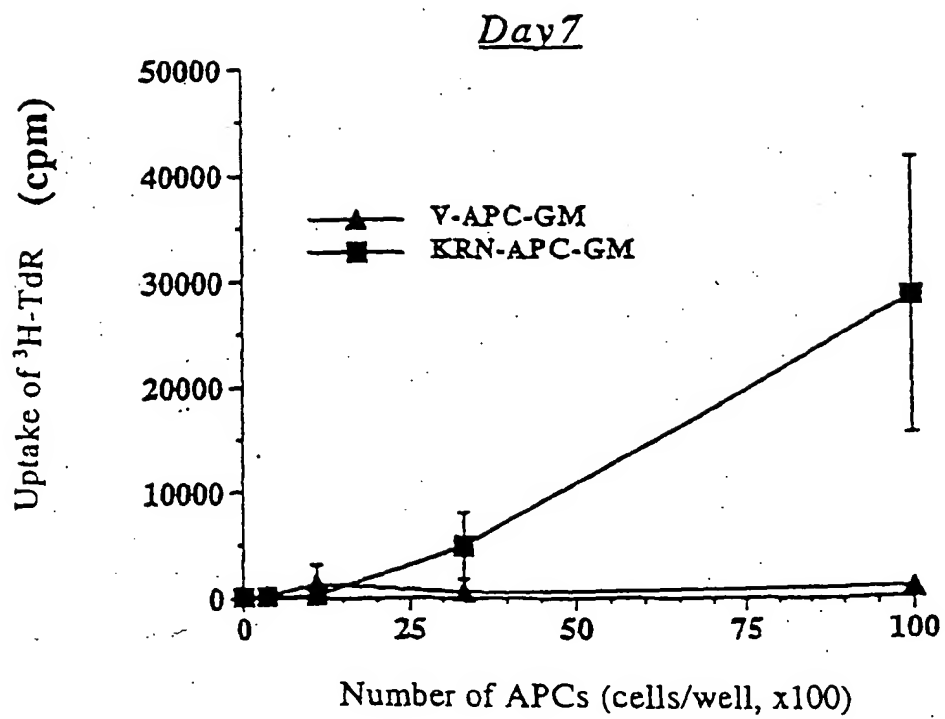


Fig. 12

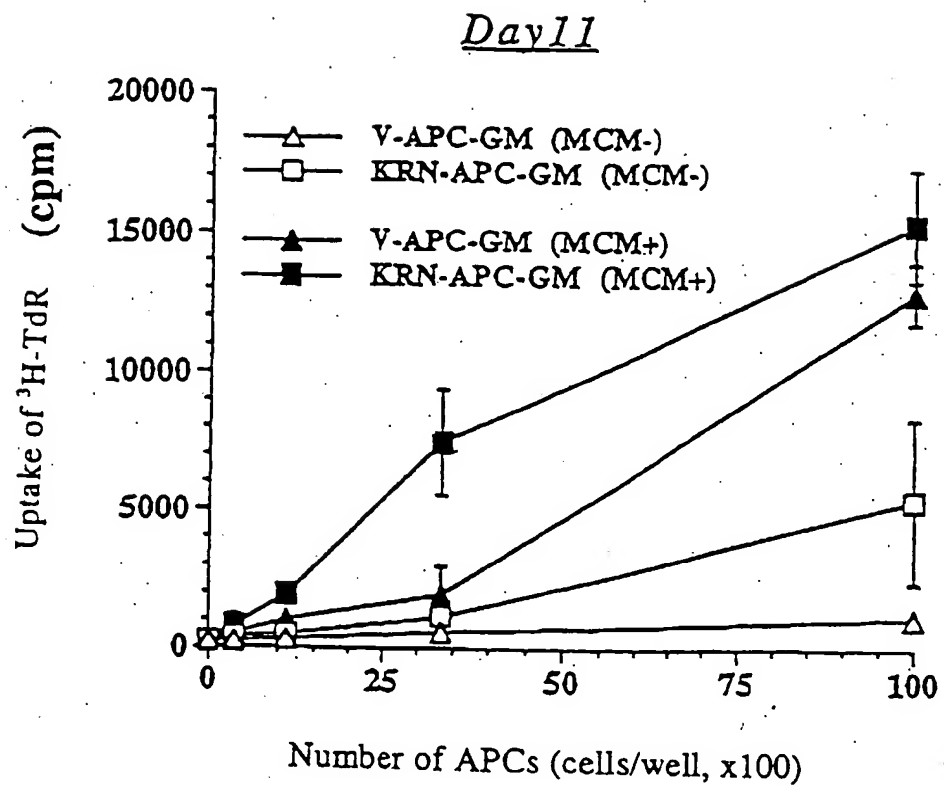


Fig. 13

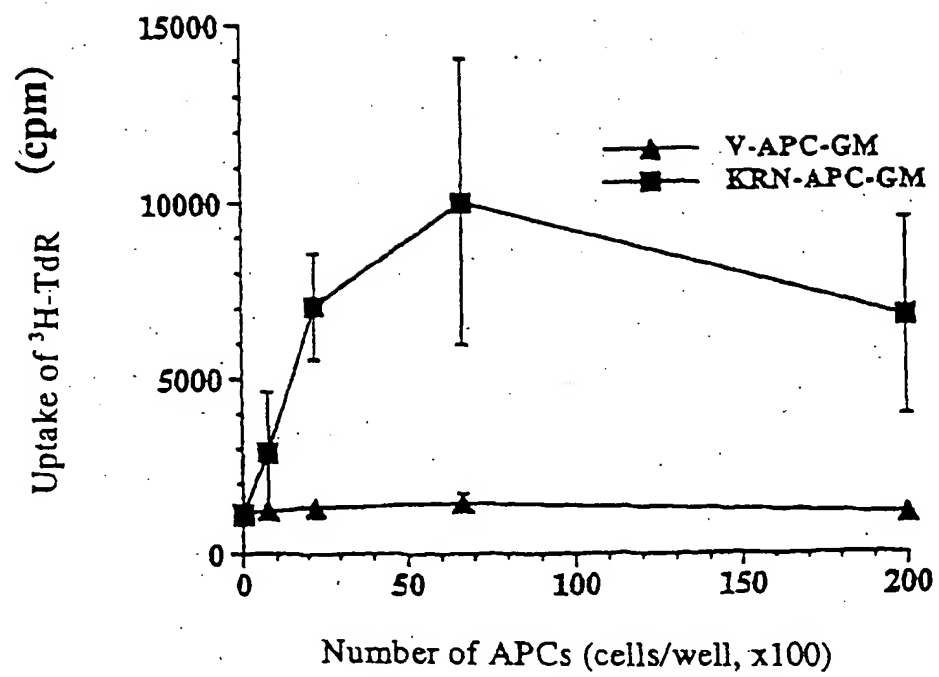
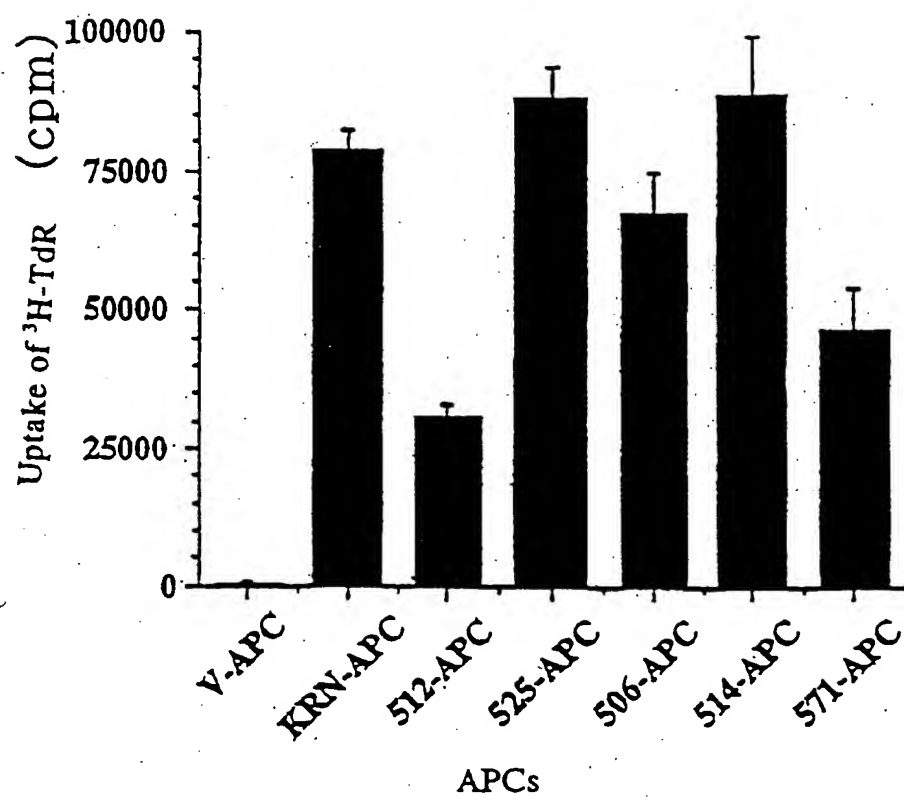


Fig. 14



INTERNATIONAL SEARCH REPORT

International application No.
PCT/JP97/04832

A. CLASSIFICATION OF SUBJECT MATTER Int. Cl ⁶ C12N5/22, C07H15/04, A61K35/14, A61K35/28, A61K35/36 // A61K31/70 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) Int. Cl ⁶ C12N5/00-5/28, C07H15/00-15/26, A61K35/00-35/84, A61K31/00-31/80 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) WPI (DIALOG), BIOSIS (DIALOG), CA (STN), REGISTRY (STN)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	YAMAGUCHI Y. et al., "Enhancing Effects of (2S,3S,4R)-1-O-(α -D-Galactopyranosyl)-2-(N-Hexacosanoylamino)-1,3,4-Octadecanetriol (KRN7000) on Antigen-Presenting Function of Antigen-Presenting Cells and Antimetastatic Activity of KRN7000-Pretreated Antigen-Presenting Cells", Oncology Research, Vol. 8 No. 10/11 (1996) p.399-407	1-10, 14-21, 23
Y	JP, 5-59081, A (Kirin Brewery Co., Ltd.), March 9, 1993 (09. 03. 93) & JP, 4-508706, A & WO, 93/05055, A1 & EP, 609437, A1	1-10, 14, 20
Y	KOEZUKA Y. et al., "Treatment of hepatic metastasis by KRN7000(II): Its immunotherapeutic mechanism", Proceedings of American Association for Cancer Research, Vol. 37 (March 1996) p.447	1-10, 14, 20
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search March 25, 1998 (25. 03. 98)		Date of mailing of the international search report April 7, 1998 (07. 04. 98)
Name and mailing address of the ISA/ Japanese Patent Office		Authorized officer
Facsimile No.		Telephone No.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP97/04832

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JP, 6-145189, A (The Kitasato Institute and another), May 24, 1994 (24. 05. 94) & JP, 4-503301, A & WO, 92/12986, A1 & EP, 639580, A1 & US 5672693, A	1-21, 23

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